ofsoilc0

7

66)

UC-NRLF

B 2 912 815

LIBRARY

OF THE

University of California.

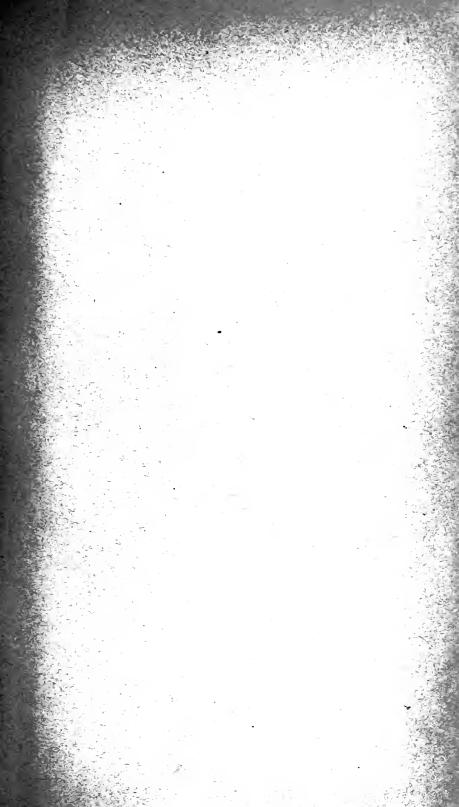
GIFT OF

usac

Class







Digitized by the Internet Archive in 2007 with funding from Microsoft Corporation

19 19 1908

U. S. DEPARTMENT OF AGRICULTURE,

BUREAU OF CHEMISTRY—BULLETIN NO. 76.

H. W. WILEY, Chief of Bureau.

INSECTICIDE STUDIES.

- I. PYRETHRUM POWDERS CONTAINING POISONOUS METALS.
- II. A COMPILATION OF ANALYSES OF INSECTICIDES AND FUNGICIDES.
- III. STATE LAWS GOVERNING THE COMPOSITION AND SALE OF INSECTICIDES.

BY

J. K. HAYWOOD,

Chief, Insecticide and Agricultural Water Laboratory.



WASHINGTON:
GOVERNMENT PRINTING OFFICE





U. S. DEPARTMENT OF AGRICULTURE,

BUREAU OF CHEMISTRY—BULLETIN NO. 76.

H. W. WILEY, Chief of Bureau.

INSECTICIDE STUDIES.

- I. PYRETHRUM POWDERS CONTAINING POISONOUS METALS.
- II. A COMPILATION OF ANALYSES OF INSECTICIDES AND FUNGICIDES.
- III. STATE LAWS GOVERNING THE COMPOSITION . AND SALE OF INSECTICIDES.

 $\mathbf{B}\mathbf{Y}$

J. K. HAYWOOD,

Chief, Insecticide and Agricultural Water Laboratory.



WASHINGTON:
GOVERNMENT PRINTING OFFICE.
1903.



LETTER OF TRANSMITTAL.

U. S. DEPARTMENT OF AGRICULTURE.

BUREAU OF CHEMISTRY,

Washington, D. C., March 15, 1903.

SIR: I have the honor to transmit for your approval three papers on insecticides prepared in the Insecticide and Agricultural Water Laboratory of this Bureau, and to recommend their publication as Bulletin No. 76 of the Bureau of Chemistry.

Respectfully,

H. W. WILEY,

Chief.

Hon. James Wilson,

Secretary of Agriculture.



CONTENTS.

P	yrethrum powders containing poisonous metals
	Introduction
	Cases of poisoning by lead and other chrontates
	Lead poisoning caused by the manufacture of fuses for lighting purposes.
	Fatal poisoning by eating confectionery colored with lead chro-
	mate
	Memoirs concerning dangers to which workmen in chromate factories are exposed
	Analyses of pyrethrum powders
	Collection of samples
	Methods and results of analysis
	Correspondence with manufacturers
Ι.	A compilation of analyses of insecticides and fungicides
	Introduction
	Sources of analyses and description of samples.
	Paris greens
	Methods
	Comment on analyses
	London, English, and Paris purples
	Insecticides and fungicides other than Paris Green and London Pur-
	ple that contain arsenic or copper, or both as active constituents.
	White arsenoids.
	Paris Green and Bordeaux Mixture
	Slug Shots
	Peroxid of silicates
	Black Death
	English Bug Compound
	Oriental Fertilizer and Bug Destroyer.
	Lead Arsenate and Pink Arsenoid
	XX, Purple Poison, and Insecticide
	Copper carbonate
	Fostite
	Sulphatine and Death to Rose Bugs
	Soaps
	Lyes
	Hellebores
	Mixtures containing borax. Hooker's Cockroach and Water Bug Exterminator
	Tobacco extracts
	Formalins
	I UIIIIIIII

II. A compilation of analyses of insecticides and fungicides—Continued.	Page.
Sources of analyses and description of samples—Continued.	
Petroleums	
Miscellaneous solid insecticides and fungicides	
Sulphurs	
Pine-tar insecticides	. 50
Bug Deaths.	. 50
Potassium cyanid	
Nonpoisonous Potato Bug Destroyer	. 51
Nonpoisonous Insect Exterminator	. 51
X. O. Dust	. 51
Electric Vermin Exterminator	. 52
Par'Oidium	. 52
Antinonnin	. 52
Miscellaneous liquid insecticides and fungicides	. 52
Bollene Insecticide	. 52
Excelsior Insecticide and Wood Preserver, Red Cedar brand.	
Fly Preventative	
Karbo-Kresolate	. 55
Professor De Graff's Carpet Bug Destroyer	. 56
III. State laws governing the composition and sale of insecticides	. 57
California	. 57
Louisiana	. 58
New York	. 59
Oregon	. 61
Texas	
Washington	. 63

INSECTICIDE STUDIES.

I. PYRETHRUM POWDERS CONTAINING POISONOUS METALS.

INTRODUCTION.

In the course of a recent examination of a large number of miscellaneous insecticides it was observed that a number of pyrethrum powders contained lead chromate. These powders are sold under such names as Persian Insect Powder, Dalmatian Insect Powder, Buhach, and Pyrethrum, and are colored yellow because there seems to be a popular demand for this grade of goods, the public apparently believing that the preparation must be bright yellow in order to kill insects. As a matter of fact pure pyrethrum, which is composed of the ground flower heads of Pyrethrum roseum and Pyrethrum cinerariæfolium, usually possesses a light yellow color, but never the bright, intense yellow color given to it by lead chromate.

As lead chromate is known to be poisonous, it would appear that it is a very reprehensible practice to add it to pyrethrum powders, since they are often blown about living rooms with a bellows to kill insects, and are consequently inhaled by human beings. Besides this, pyrethrum is always advertised to be harmless to human beings, which it is when pure, but ceases to be so when it has been mixed with lead chromate or other yellow colored chromates. In view of the above facts it was deemed advisable to examine the pyrethrum powders sold upon the American market to discover which of them contain yellow chromates and at the same time to cite cases of poisoning by chromates to substantiate the claim that any of the yellow chromates that might be used for this purpose, and especially lead chromate, are poisonous to human beings. The facts which have been collected on these points are embodied in the following report:

CASES OF POISONING BY LEAD AND OTHER CHROMATES.

Since it is not generally known that many pyrethrum powders contain lead chromate, it doubtless often happens that physicians are much puzzled as to the cause of the illness of persons who have been exposed to the action of such powders. In order, therefore, that physicians and others may recognize the symptoms of chromate

poisoning caused by this class of goods, the following account is given of cases of chromate poisoning found in medical records:

LEAD POISONING CAUSED BY THE MANUFACTURE OF FUSES FOR LIGHTING PURPOSES, a

On February 22, 1875, a woman of about 46 years of age came under my care at the hospital. She complained of pains in the stomach and a general indisposition or breaking down. For a number of days she had suffered with severe gripes. Her stomach was not swollen, but her skin was very sensitive and palpation caused a contraction of the abdominal muscles. There was, however, no tumor in the abdominal cavity. She had suffered for about fifteen days with obstinate constipation that could be relieved only by an injection; her liver was small and not painful; her tongue white, appetite nearly entirely lacking, and breath foul. There was nausea and a distaste for food, and one could observe the existence of a grayish-black slate-colored border on the edges of her teeth next to the gums, which had all the characteristics of lead bands. That symptom, added to the gripes and to cramps in the calves of the legs during the night, caused me to diagnose the case as lead poisoning. This woman was under my care until the 16th of March, at which time she began to convalesce, although at that time she was still anæmic and feeble; but there was unquestionable improvement.

Being well convinced that my patient was suffering with lead poisoning, I thought I ought to find the cause. I found that she had worked for five years in a well-aired factory employing about 30 people, where they were engaged in the manufacture of fuses which were to be used by smokers to light their pipes and cigars. Her work differed from that of her fellow-workers in that she was employed in winding the cotton that was to be used for the manufacture of fuses, while they wove the threads of cotton. She was thus more exposed than her companions to the emanating dust, especially as the cold weather had obliged them to close the windows of the shop. These fuses were about as large around as a pencil or the little finger and of a yellowish orange color, indicating that they were colored with lead chromate. A chemical analysis showed the presence of this compound.

At the shop where these fuses were manufactured I found that those who were engaged in weaving them did not show any appreciable symptoms of lead poisoning, while those who were employed to shake out and wind the cotton all had fairly well developed cases. The merchant at Paris who manufactured these fuses said that they were originated in Germany, but were now exported largely, especially to America.

FATAL POISONING BY EATING CONFECTIONERY COLORED WITH LEAD CHROMATE. h

On March 14, 187-, between 9 and 11 a. m., the two sons of Mr. H., aged respectively 3½ and 1¾ years, ate several small ornaments representing bees that had been placed on a cake. In the afternoon of the same day they became very ill and were attacked with severe vomiting and great prostration. The vomited mass was at first yellow, and was thrown up with great pain. Vomiting continued until 11 p. m., after which it became infrequent. When the physician arrived, at 6 p. m., they were in bed and suffering extremely. They had reddened faces, complained of great thirst, were languid, had no dysentery, although they complained of slight pains.

a Annales d'hygiène, 2d series, No. 44, 1875.

^b Vierteljahresschrift für gerichtl. Medizin, 1874, 20: 60.

It was easy to see that they had been poisoned, and since in the morning they had eaten the same food as the rest of the household, who remained in perfect health, probably the poisoning was due to the above-mentioned bees, seven of which had been given to the children to play with, while only one could be found. They finally confessed that they had eaten the remaining six, and the nurse had seen them do it. The nurse thinking it might be harmful, had taken a softened "bee" from the younger child and wiped out his mouth.

One of the "bees," of which more were found at the confectioner's who had made the cake, was examined by a chemist and found to consist of gum and lead chromate. On the following day both patients had the same red and heated faces; they showed no interest, and on being questioned gave tokens of displeasure. They did not complain of local pains, evidently because their senses were benumbed. The younger had some diarrhea and toward noon convulsions, from which his face was livid. The convulsions became more frequent toward evening, and at 9 o'clock on the morning of the following day, March 16, the younger boy died.

The older child still had a red, heated face and displayed no interest, being almost unconscious. The skin upon his breast and abdomen was strikingly red. The evening temperature was 39.5° in the armpits. On the 17th the pulse was irregular and the temperature in the evening 39.6°. Drinking was extremely painful, in spite of great thirst. On the 18th a foul smell came from the mouth, redness was still present, accompanied by stupidity and painful swallowing; temperature in the evening was 39.2°. On the 19th a collapse took place, a very foul smell came from the mouth, and swallowing was almost impossible. The senses were very much benumbed, and death followed at 11 a. m. The result of the post-mortem may be summed up in tle words of the physician as follows: "The death of the children is the direct result of the destruction which has befallen the digestive canal from the throat to the duodenum. Concerning the cause of this destruction and concerning some further conditions we will not express an explicit opinion." The liver, stomach and contents, and urine were then subjected to a chemical examination. This gave negative results, except that in the liver of both corpses traces of copper were found, a discovery of no toxological interest. The absence of all poisonous metals in all other parts of the corpses was shown.

The "bees" were also examined, and 0.27 gram of gum and 0.0042 gram of neutral lead chromate were found in one that was 13 mm long and 5 mm thick. The powder used by the confectioner was also analyzed and found to contain lead chromate. The illness of both children, occurring at the same time and showing itself by severe vomiting, must be traced back to their eating some poisonous food; and since it was known that they both ate the "bees" which contained lead chromate and that they vomited up yellow vomit, the eating of the metallic salt must be considered as the cause of the deaths.

It is not remarkable that no lead chromate was found in the alimentary tract, since the children vomited a great deal and thus threw up all of the poison.

FATAL POISONING BY INHALING DUST FROM YARN COLORED WITH LEAD CHROMATE. $^{\prime\prime}$

On December 29, 1876, a weaver, B., at E., received a package of yellow threads, colored with lead chromate, which were to be used for the manufacture of curtains. B. and his mate, G., worked at one spool and Mrs. B. and a girl, L., at the other. They worked from 7 a. m. to 10 and 11 p. m. While they were working the threads shed powder, according to B.'s statement, so that their faces and hair became entirely

yellow. Their spittle appeared yellow, and the powder had a bitter taste. According to Mrs. B. the powder had a sweetish-bitter taste. The male helper sickened after working upon the threads about eight days; he had a headache, with a ringing in the ears, pains in his breast and stomach, and lost his appetite. He had a desire to vomit and was constipated. This continued for six weeks, until he ceased working and went to the doctor. Mrs. B. during the meantime became ill; she complained of loss of appetite and pains in the chest. B. on the last day of the third week of the work was attacked with pains in the stomach. Then appeared nausea, languor, and sleeplessness. Eight or ten days later he was obliged to stop work and send for the doctor on account of pains in the bowels and costiveness. The girl helper was also ill, having diarrhea, loss of appetite, and pains in the breast.

Dr. J. stated the symptoms of B. and the helpers in the following terms: Yellow-coated tongue and yellow expectoration, entire loss of appetite, nausea, temporary vomiting, and pains in the region of the navel, accompanied by obstinate constipation. The excrement was also colored yellow, and all the patients were very weak. The doctor stated that these symptoms were all caused by breathing yellow dust from

the yarn and indicated plainly lead-chromate poisoning.

Although all of the above-mentioned persons finally recovered, B. lost a 9-weeksold baby, whose death could be ascribed only to the breathing or swallowing of the varn dust. When the B.'s noticed how much dust was flying about they sought to protect their baby by spreading a white woolen cloth of medium thickness over his face. This cloth was said to have been vellow with the dust all the time, as well as the bed on which the child was lying. The child's crib stood midway between the two B.'s stools and at night the mother took the child to bed with her. healthy and lively, so that continued poor feeding could not have caused the bad condition of the stomach (jelly-like softening and perforation) observed at the postmortem. Suddenly, six or eight days before its death, which occurred on February 24, the child was taken sick. Its face became pale and its body hot. It had a couple of loose, yellow actions each day, which were accompanied by restlessness and frequent screaming. The skin upon the breast and stomach became red. the beginning the baby drank, but refused to eat; later it drank with trouble, and on the day it died it swallowed with difficulty. The lips were dry, the breathing short, and death followed slowly. It is to be remarked that the child, after remaining healthy for six weeks in spite of the surrounding yarn dust, was suddenly taken sick six or eight days before death and grew worse at a time when the yellow yarn was not being worked upon. Since the child remained covered with the cloth up to the time of its death, it appears likely that the dust gradually worked its way through the cloth and sifted down so that the child always had to breathe this impure air until its death.

 Λ chemical examination was made of the following organs of the child and the articles that were used at the the time of his death: Contents of the nose, contents of the stomach, urine and gall taken from the urine and gall bladders, organs of the stomach and breast, the hair of the head, shirt slip, the child's nipple, and a portion of the yellow yarn. The brain would have been examined, but the vessel containing it was lost. The yarn contained 11.83 per cent lead chromate. Lead chromate was found in the hair of the head, and the little slip contained 65 mg lightly strewn over it. The organs of the neck, chest, and gullet contained 36 mg of lead chromate. The other fluids of the body and organs, the shirt, and nipple did not show the presence of any lead chromate.

It will thus be seen that numbers of authentic cases of poisoning, by either breathing or swallowing lead chromate, are on record. Following will be found several typical cases of poisoning by alkaline chromates and bichromates selected from the large number that have been observed by Messrs. Delpech and Hillaire: a

MEMOIRS CONCERNING DANGERS TO WHICH WORKMEN IN CHROMATE FACTORIES ARE EXPOSED.

Observation I.— Workmen employed for a long time at the grinder (pilerie) and formerly at the kettles (chandières)—Nasal partition perforated by the dust.

V., aged 50, has worked for twelve years (1849) in a chromate manufacturing plant. He was employed from the beginning at the grinder and sifter, and is thus constantly enveloped in an atmosphere charged with fine chrome-iron powder. He says that he feels no indisposition. He wipes his nose and spits blackish matter, but neither coughs nor has any smothered feeling; his health appears to be good; he has no disease of the lungs; he does not believe there is anything the matter with his nose; however, his sense of smell was perfect when he entered the factory, and he has now completely lost it. The nasal passages were examined, and a perforation of the cartilage was observed. He says that since entering the factory he has sometimes worked at the kettles. At the commencement of this labor he was attacked with headache and swelling of the nose accompanied by an abundant discharge from the nostrils. His eyes did not smart. These symptoms lasted about fifteen days, after which he regained the most perfect health without even experiencing again the symptoms enumerated, although employed at the same work.

This workman has since (October, 1864) died after a few days' illness of inflammation of the chest, without having previously shown any prolonged symptoms of suffering with the respiratory organs.

Observation III.—Workman employed at the kettles for about four weeks—Inflammation of the mucous membrane of the nose—Sore upon the right thumb—A nasal perforation—Ulceration of both of the external nasal walls.

E., aged 28, who had a good constitution, had been employed for five weeks in a potassium bichromate manufacturing plant when he was examined by us. He was employed about eight days inside the works carrying the bichromate and for three weeks at the kettles. From the first day he was attacked by prolonged and intense sneezing. After four days he suffered greatly with intense prickling, abundant running of the nose, an incessant desire to blow his nose, and the passage of red membraneous shreds. He had no watering of the eyes, or at most it was slight. The most severe symptoms lasted from ten to twelve days. He had nearly recovered when he stopped working at the kettle, and has since completely recovered. This workman still has his sense of smell. He was afflicted by a sore on his right hand, caused by penetration of particles of bichromate in a slight excoriation, and a slight ulceration still remains. One can also note brown spots, caused by sores of the same nature, upon the side of his face and on the ring finger.

E. has never suffered with his feet. He has no eruption either on the skin of his body or genital organs. His eyelids are not excoriated, and he has neither vomited, suffered with diarrhea, nor a cough. The nasal partition is perforated at the usual place.

a Annales d'Hygiène Publique et de Medicine Legale, 1869, and 1876, 2d series.

Observation VI.—Workman, a coppersmith unaccustomed to the manufacture of chromate, was exposed to the dust resulting from the repairing of the kettles—Inflammation of the mucous membrane of the nose and nasal perforation.

F., a coppersmith and a stranger to the chromate factory, came into the factory for a short time to repair the kettles. This repairing required a readjusting of the sides by hitting with a hammer, and these blows caused the crust of bichromate that adheres in the slightest irregularities to fly about in the form of a dust. He was attacked quickly by inflammation of the mucous membrane of the nose and lost his nasal partition in eight days.

Observation XI.—Superintendent, not employed at any special work—Chronic irritation, swelling of the membrane at the usual place, but no perforation.

O., aged 20 (December, 1863), has been employed for four years in the chromate works. He has never worked at the kettles or the calciner. Charged with general surveillance, he merely walks through the works; in exceptional cases, however, he helps solder the lead crystallizing vats that are in need of repairs. He has consequently to a certain extent been subjected to the action of bichromate. At the time we examined him he had felt a tingling of the nose and had frequent fits of sneezing, but his nasal membrane had not been badly inflamed. His nasal partition is not perforated, but at the usual point he has a slight swelling and reddening of the membrane. His general health is good.

Observation XVIII.—Fireman exposed to chromate dust—Perforation of the nasal cartilage, ulcerated sores on the right hand.

D., aged 28, has been employed at the works for two years, and for the last ten months at the manufacture of potassium chromate. He is employed as a fireman, but often helps his fellow-workmen, and so during much of the time spent in the factory he is exposed to chromate dust. He had always enjoyed the best of health. He had never had syphilitic symptoms, but during his infancy the glands at the side of his neck were swollen for a long time. His general condition is good. He had worked at the manufacture of potassium chromate for scarcely fifteen days when he was taken with nose bleeding, sneezing, pain in the nose, and watering of the eves. He sometimes experienced a little uneasiness in breathing. It was fifteen or twenty days after he had ceased to work at the preparation of potassium chromate that he examined his nose and found a small perforation. The nose preserves its normal form. There is an ulcer on the nasal cartilage in the right nostril, and at the edges of this the mucous membrane is very much inflamed. In the left nostril is an opening corresponding to this ulceration. His sense of smell is preserved; neither the throat, palate, nor roof of the mouth have been attacked, and he has no cough. This workman has hard, cartilaginous, livid sears upon his thumb and his right hand. On the left index finger there exists a sore which is much swollen and exudes a sero-purulent liquid. This workman has never had any swelling or ulceration of his feet.

There are many other observations cited in the above article, but enough have been given to show that persons who are obliged to breathe air charged with the dust from chromates or bichromates are nearly always affected more or less seriously.

ANALYSES OF PYRETHRUM POWDERS.

COLLECTION OF SAMPLES.

In order that samples of pyrethrum, representative of all grades, might be obtained, a circular letter was forwarded to the chemist of each of the experiment stations, with the request that they purchase for us, upon the open market, three or four samples of those products sold under the name of either Pyrethrum, Buhach, Persian Insect Powder, or Dalmatian Insect Powder. They were also asked to give data as to the price per two-ounce sample, the name and address of the retail dealer, and the name and address of the person who originally prepared the sample. In answer to these letters 105 samples were received, all of which were subjected to chemical examination. It is believed that these samples represent fairly well the present state of the American market.

Credit is due to the assistants in the laboratory, Bernard H. Smith and Harry W. Houghton, who aided in performing the analytical work.

METHODS AND RESULTS OF ANALYSIS.

Since lead, potassium, and barium chromate have all a bright yellow color, and any of them might be used to color the pyrethrum powders, tests for all of these substances were made in the samples under examination according to the following methods:

Qualitative test for lead.—Ash about 2 grams of the sample in porcelain and treat with a little water and about 20 cc of concentrated hydrochloric acid in a porcelain dish. Place on the steam bath and steam for about one-half an hour or until all the lead is in solution and the chromium reduced and in solution. Filter after dilution. Add sodium carbonate until the solution is alkaline and then add acetic acid until the solution is just acid. Filter and add a solution of potassium chromate. A vellow precipitate or cloudiness indicates lead.

Qualitative test for barium.—Ash about 2 grams of the sample in porcelain and treat with hydrochloric acid as directed for lead. Evaporate to dryness and take up with about 0.8 ec of hydrochloric acid. Filter and pass hydrogen sulphid through the filtrate. Filter off any sulphids that may precipitate and test for barium in the filtrate with a few drops of dilute sulphuric acid.

Qualitative test for chromium.—Ash about 2 grams of the sample in porcelain. Treat this with about four times its weight of a mixture of equal parts of sodium carbonate and potassium nitrate, and fuse. Dissolve in water on the steam bath and filter. Make slightly acid with acetic acid and test with lead acetate for the presence of chromium.

Testing the powders by the above methods showed that whenever chromium was present lead was also present, these two evidently being combined as lead chromate. A quantitative estimation was also made of the chromium present according to the following method of analysis and the chromium was calculated to lead chromate.

Table I.—Description of sumples, source, and price.

		Netarici.	Mannfacturer or wholesaler.	2 ounces.
98 B	Persian Insect Powder Dalhadian Insect Powder	G. B. Evans, Philadelphia, Pa. Llewellyn, Philadelphia, Pa.	G. B. Evans, Philadelphia, Pa. Llewellyn, Philadelphia, Pa	 9.00 1.00 1.00 1.00 1.00 1.00 1.00
	Pyrethrum	Menges, Madison, Wis	Allaire, Woodward & Co., Peoria, III	: :
0.75	op	Henry Lewis, Madison, Wis	Fuller & Fuller, Chicago, Ill.	-
E 25	Dalmathan Insect Powder	Hollister's Pharmacy, Madison, Wis	Alhaire, Woodward & Co., Peoria, III	-
	balmatian Insect Powder	Robinson Drug Co. Lansing, Mich	do	
-	Spalato Insect Powder	Baner's Drug Co., Lansing, Mich.	Lehn & Fink, New York, N. Y.	
	Palmatian Insect Powder	M. A. Diekinson, Amherst, Mass	Archibald & Lewis, New York, N. Y	
	Persian Insect Powder	W. S. Wescott, Amherst, Mass.	Hoyt Medicine Co., New York, N. Y	-
	Dalmatian Insect Powder	Holland & Gallord, Amberst, Mass	Lee & Osgood, Norwich, Conn.	•
200	40	H. Adams, Amberst, Mass.	E. L. Patch Co. (wholesaler), Boston, Mass.	:
_	disect Fowder	Doubles Denel, Amberst, Mass.	Alloing Windmand E. Co., Decailed 111	-
	do	Harlot Drug Co. Lincoln Nobe	Americ, woodward & Co., Feoria, III	:
	Subach	H. W. Brown, Lincoln Nebr	Ruhach Manufacturing Co. Stockton, Cal	:
	vrethrum		Richardson Drug Co., Onatha, Nebr.	:
_	nsect Powder	C. Powells, Stillwater, Okla		
	Pyrethrum	W. H. Hand, Stillwater, Okla	Parke, Davis & Co., Detroit, Mich	
_	Insect Powder	i		:
	'yrethrum	Koller & Masters, Manhattan, Kans		:
	ор	G. B. Harrot, Manhattan, Kans	Faxon, Horton & Gallagher, Kansas City, Mo	:
	minimitian Insect Fowder	Orr Drug Co., Athens, Ga	(filpin, Langdon & Co., Baltimore, Md	:
	00	Falmer & Kumebam, Athens, Ga		:
	Persum Insect Fowder	H. O. Meek, State College, Pa	Smith, Kine & French, Philadelphia, Pa	:
255	Suhach	George Martin, Tueson, Ariz		:
	Tyrethirm	D. L. Kead, bryan, 1ex	Firke, Davis & Co., Detroit, Mich	:
-		M. H. T. L. Dans Co. Co.	_	:
375	Persian Insect Powder	The Van Densen Pharmacy, New Brunswick.	Mckells Stone-Chemical Co., New Orleans, Ed.	: :
		N. J.		
	op	William Rust & Sons, New Branswick, N. J	William Rust & Sons, New Brunswick, N. J	:
		Skillman & Van Pelt, New Brunswick, N. J		
-	Insect Powder	L. Christianson, Fargo, N. Dak		:
	dp	R. H. Finney, Fargo, N. Dak		:
	op	Font & Portfield, Fargo, N. Dak	_	-
	op	B. F. Mackaff, Moorhead, Minn		-
-	op	Waldorf Pharmacy, Fargo, N. Dak	Parke, Davis & Co., Detro t, Mich	:
٠		G. C. Sadin, Fargo, N. Duk		:
-	Pyrethrum	J. E. Knight, Exeter, N. H		-
•	(do	II		
000	Description Institute Description	J. C. Vickery, Dover, N. H.		:
	reising insect rowder	II. E. CHEK, Lattalyette, Ind		:

PYRETHRUM POWDERS CONTAINING POISONOUS METALS.	13
8:252556 8:3526 12256	07.
Stolhman & Fulton, New York, N. Y F. H. Leggett, New York, N. Y Schleffelin, & Co., Detroit, Meh Parke, Davis & Co., Detroit, Meh Merkeson & Robbins, New York, N. Y Burlington Drug Co., Bration, Net Burlington Drug Co., Bratington, Vt Parke, Davis & Co., Detroit, Mich Alhaire, Woodward & Co., Peorin, Ill Alhaire, Woodward & Co., Peorin, Ill Butharch Manufacturing Co., Stockton, Cal Lehn & Fink, New York, N. Y Parke, Davis & Co., Detroit, Mich do. Leggett Boxe, Boston, Mass Nathan Howe, Boston, Mass B. Ritchey & Co., New York, N. Y B. Ritchey & Co., New York, N. Y B. Ritchey & Co., New York, N. Y Bullarch Manufacturing Co., Stockton, Cal Albaire, Woodward & Co., Peorin, Ill Parke, Davis & Co., Detroit, Mich Bullarch Manufacturing Co., Stockton, Cal Albaire, Woodward & Co., Peorin, Ill Parke, Davis & Co., Detroit, Mich Albaire, Woodward & Co., Peorin, Ill Parke, Bavis & Co., Detroit, Mich Albaire, Woodward & Co., Peorin, Ill Parke, Bavis & Co., Detroit, Mich Albaire, Woodward & Co., Peorin, Ill Parke, Bavis & Co., Peorin, Ill Parke, Novel Verk, N. Y.	Parke, Davis & Co., Petroit, Mich.
Bogan-Johnson Drug Co., Lafayette, Ind. B. Sloam, Chemson College, S. C. E. G. Farnis, J. A. Co., Pendleton, S. C. C. A. Barber, Burlington, Vt. G. A. Churchhill, Burlington, Vt. G. E. Perkins Burlington, Vt. G. E. Perkins Burlington, Vt. T. Bellerbab, Burlington, Vt. Crystal Pharmary, Burlington, Vt. Graham & Wortham, Corvallis, Oreg Graham & Wortham, Corvallis, Oreg Graham & Wells, Corvallis, Oreg Graham & Wells, Corvallis, Oreg Owens & Owens, Wytherville, Va. While & Burdiek, Hunea, N. Y. Brooks's Drug Store, Hartford, Conn. A. Dutschheyer, Hartford, Conn. C. I. Belle, Hartford, Conn. C. L. Finel, Norwalk, Con	C. T. Carr & Bro., Hyattsville, Md Thomson & Hogan, Christiansburg, Va
Main	
2.8.8.8.8.8.8.8.8.8.8.8.8.8.8.8.8.8.8.8	657

Table 1.—Description of samples, source, and price—Continued.

Price per 2 onnees.	98 - 2.2 - 10 - 10 - 10 - 10 - 10 - 10 - 10 - 10
Manufacturer or wholesaler.	R. S. Chrson, East Bradford, Va. Parks, Davis & Co., Detroit, Mich. 80, 23, 43, 44, 45, 46, 46, 47, 47, 47, 47, 47, 47, 47, 47, 47, 47
Retailer.	R. S. Curson, East Bradford, Yu. G. W. Lyke, Bradford, Va. C. L. McCharlor, Moscow, Idaho A. K. Greenman, Kingston, R. I. Z. D. Gilman, Washington, D. C.
Name,	H. S. Carson, Bast Bradford, Yn. 649 Fersian Insect Powder C. W. Lyle, Bradford, Au. 640 Haver Powder C. W. Lyle, Bradford, Au. 641 Haver Powder C. M. Carson, Markow, Idaho C. M. Greenman, Kingston, R. I. H. Torsen, Moscow, Idaho A. K. Greenman, Kingston, R. I. Greenman, R. I. Greenman
Xerial No.	655 655 655 655 655 655 655 655 655 655

 α General index number.

Method for determining chromic acid.—Weigh out from 2 to 4 grams of the sample. Ash in a porcelain dish and allow to cool. a little water and a large amount of concentrated hydrochloric acid and · evaporate to dryness. Treat again with concentrated hydrochloric acid and evaporate to dryness.

Take up the residue with about 0.8 cc of concentrated hydrochloric acid with the aid of heat. Filter and pass hydrogen sulphid through the solution until all lead is precipitated. Filter and evaporate the filtrate to dryness. Take up the residue with a little water and 1 cc of hydrochloric acid with the aid of heat. Precipitate out the chromium (with iron, aluminum, and some phosphoric acid) by means of ammonia, taking care not to add any excess of ammonia. Filter and wash with hot water. Dissolve the precipitate on the filter by means of hot dilute hydrochloric acid and evaporate to dryness. Take up with a little hot water and 6 to 8 drops of concentrated hydrochloric acid, heating for a short time if necessary. Transfer to a 50 ce or 100 ce flash and make up to the mark. Now prepare an alkaline permanganate solution in the following manner: Make an approximately N 10 solution of potassium permanganate and standardize it against ammonium ferrous sulphate. Measure out exactly a liter of this solution and add a few grams of sodium carbonate and 1 gram of sodium hydroxid and make the entire volume up to 2 liters. This is the standard solution required.

A convenient number of cubic centimeters of this standard is measured into a beaker and about 50 cc of distilled water, accompanied by a little sodium carbonate, is added. The solution of chromium previously prepared is then run in from a burette, while the contents of the beaker are boiling, until the pink color is dissipated and a pure vellow color results. Knowing the strength of the permanganate and the number of cubic centimeters of the chromium solution used, one can calculate the percentage of chromium in the original powder according to the following equation:

$$\mathrm{CrCl_3} + \mathrm{KMnO_4} + 4 \ \mathrm{KOH} = \mathrm{K_2CrO_4} + \mathrm{MnO_2} + 3 \ \mathrm{KCl} + 2\mathrm{H_2O}$$

In Table II are found the results obtained upon the samples of pyrethrum powder examined:

Table II.—Examination of pyrethrum powders for coloring matter.a

Serial No.	Name.	Chromium.	Lend.	trioxid	Lead chro- mate (PbCrO ₄).
266 267 268 269 270 271 272		Present do do do	I'resentdoAbsentdododododododo	0.099 .071	0.32
	a Examination for barium gave no	cative result	s for all san	inles	

Table II.—Examination of pyrethrum powders for coloring matter—Continued.

Serial No.	Name.	Chromium.	Lead.	Chromium trioxid (CrO ₃).	Lead ehro- mate (PbCrO ₄).
		_		Per cent.	Per cent.
a 273	Dalmatian Insect Powder	Present	Present	. 068	. 22
274 275	Spalato Insect Powder	Absent Present	Absent	. 068	.42
276	Persian Insect Powder	Absent	Absent	. 130	.42
277	Dalmatian Insect Powder	Present	Present	068	. 22
278	do	Absent.	Absent .	.000	į .
279	Insect Powderdododo	do	do		
280	do	do	do		
281	do	do	do		
282	Ruboob	do	l do	I .	
283	Pyrethrum Insect Powder	do	do		
312	Insect Powder	do	do		
313	Pyrethrum	do	do		
314	Insect Powder	Drosont	Procent	104	40
315 316	Pyrethrum do Dalmatian Insect Powder.	do	rresent	109	.40
317	Delmetien Insect Powder	Absent	Absent	.102	. 00
318					
a 319	do Persian Insect Powder Buhach Pyrethrum do	Present	Present	186	. 60
323	Buhach	Absent	Absent		
324	Pyrethrum	do	do		
325	do	do	do		
326	do . Persian Insect Powder	Present	Present	. 251	.81
327	Persian Insect Powder	do	do	. 365	1.18
328	do	Absent	Absent		1.36
329	do	Present	Present	. 421	1.36
330	Insect Powder	Absent	Absent		
331 332	do	do	do		
333	do do do	do	do		
334	do	do	do		
335	do Pyrethrum do	do	do		
336	Pyrethrum	do	do		
337	do	do	do		
338	do	do	do		
339	Persian Insect Powder	do	do		
a 340	Pyrethrum	Present	Present	. 090	.29
341	do do Persian Insect Powder Pyrethrum do Dulmanian Insect Powder	Absent	Absent		
384	Dalmatian Hisect Powder	00	do	• • • • • • • • • • • • • • • • • • • •	
285 a 386	do	Dwogowa t	Decoupt	0.19	1.4
387	Ineget Powder	Tresent	Present	.045	.14
388	Persian Insect Powder	do	do		
389	Persian Insect Powder Insect Powder Persian Insect Powder do	do	do		
390	do do Insect Powder. do	do	do		
391	do	do	do		
392	Insect Powder	do	do		
393	do	do	do		
394		do	op		
395	Buhach	do	do		
396	Persian Insect Powder do Dalmatian Insect Powder	00	00		
397 401	Dalmatian Insact Paudar	do	do		
402	Pyrethrum Persian Insect Powder Black Flag Insect Powder Pyrethrum do Insect Powder Princet Powder	do	do		
403	Persian Insect Powder	do	do		
404	Black Flag Insect Powder	do	do		
419	Pyrethrum	Present	Present	.037	, 12
420	do	do	do	. 056	.18
421	Insect Powder	Absent	Absent		
422		do	do		
423	Insect Powderdo	Present	Present	. 186	.60
424	do	Absent	Absent		
425 426	do	do	do		
426	Dalmatian Insect Dandon	Dec	Dr	200	1 17
427	do . Dalmatian Insect Powder . Insect Powder .	Alwant	Absont	. 502	1.17
1:20	do	da	do		
430	do	do	do		
431		do .	do .		
432		do	do		
433	do	do	do		
43.4					
	1	do	do		
435		op			
435 492	Dalmatian Insect Powder	do	do		
435	do Dalmittiin Insect Powder Bilhach. Pyrethriin	do	do		

a See correspondence with manufacturers, p. 20.

Table II.—Examination of pyrethrum powders for coloring matter—Continued.

Serial No.	Name.	Chromium.	Lead.	Chromium trioxid (CrO ₃).	Lend chro mate (PbCrO ₄).
400	Insect pourder	Absont	theont	Per cent.	Per cent.
496 509	Insect powder	Absent	Absent	.046	. 1
510	Persian Insect Powder				
511	Pyrethrum				
512	Dalmatian Insect Powder	do	do		
513	Buhach	do	d		
591	do				
592	do				
593	Persian Insect Powder				
594	Buhach				
642	Pyrethrum				
647	Insect Powder				
648	do				
649	Persian Insect Powder				
693	Buhach				
694	Insect Powder				
695	Persian Insect Powder				
696	do				
697	Insect Powder				
19530	Persian Insect Powder				1.

a General index number.

A glance at this table shows that wherever the powders are colored by a foreign substance lead chromate is the compound used. Nineteen of the 105 samples, or 18 per cent, were colored by lead chromate, the amount of which varied from 0.12 to 1.47 per cent.

Of the adulterated samples, 5 were sold under the name of Dalmatian Insect Powder, 5 under the name of Persian Insect Powder, 7 under the name of pyrethrum, and 2 simply as insect powder. It is also evident from Tables I and II that some firms are sending out two grades of pyrethrum, one containing lead chromate and the other not containing that substance. It is unfortunate that the names of all those who originally prepared the above samples could not be obtained since the powders containing the largest amount of lead chromate lack the name of the manufacturer with surprising regularity.

It seems hardly necessary to call attention to the widespread harm that is undoubtedly caused by powders thus colored. They are sold in all parts of the country and are especially advocated to kill or drive away flies, fleas, cockroaches, and other household pests, and thus they are used principally in the home. Consequently they are breathed by many people for days and weeks at a time. Such powders not only possess the poisonous characteristics of nearly all lead salts, but in addition have the toxic properties of chromic acid. It would appear, therefore, that the influence of lead chromate day by day no doubt occasions many of the mysterious cases of lead poisoning in which it seems impossible for the physician to find the source of the lead taken into the system.

CORRESPONDENCE WITH MANUFACTURERS.

In order that the manufacturers might have an opportunity to comment upon the analyses before they were published, a circular letter

was sent to each one whose goods had been examined, giving him the results of our analysis and asking for any comments that he might desire to make. Such parts of the answers as throw any light upon the subject are given, as follows:

Serial No. 340.

* * You say that the Varoter-Boggs Drug Company, of Lafayette, Ind., furnished you with a sample of our insect powder containing 0.29 per cent of lead chromate. If such is the case, they added the lead chromate themselves, as we do not use it. We do not color our insect powder, but leave that for the customers to do if their trade demands a highly colored powder.

Allaire, Woodward & Co.

Comment: It appears that the above sample was probably colored after it left the hands of the manufacturer, since seven other samples of Allaire, Woodward & Co.'s goods (Nos. 269, 271, 280, 281, 394, 496, and 591) contained no lead chromate.

Serial No. 273.

* * From the manager of our importing department we also receive the positive assurance that the powder entering into your sample No. 273 could not possibly have contained lead chromate as it left the hands of his department. It goes without saying that we never add lead chromate or any other foreign substance to our insect powder and that the adulteration which you report must have been deliberately put into it by some interested person after it passed out of our possession.

Parke, Davis & Co.

Comment: It appears from investigations which have been made that the above sample received the lead chromate after it left the hands of the manufacturer named, if indeed it were manufactured by the firm originally. Fourteen other samples of Parke, Davis & Co.'s goods (Nos. 272, 313, 331, 334, 385, 391, 402, 403, 510, 511, 647, 648, 649, 694) contained no lead chromate.

Serial No. 386.

* * * Messrs. R. D. Stearns & Co., Burlington, Vt., are in error as to the source of the Persian insect powder which they furnished your representative, stating that it came from our firm, and we have written them to that effect and asked for an explanation. We have examined their entire account for the year 1902, and positively state that they did not purchase any insect powder of us, and if they had it would have been found to be perfectly pure, as we do not allow any adulteration of Persian powder sold by us.

Schieffelin & Co.

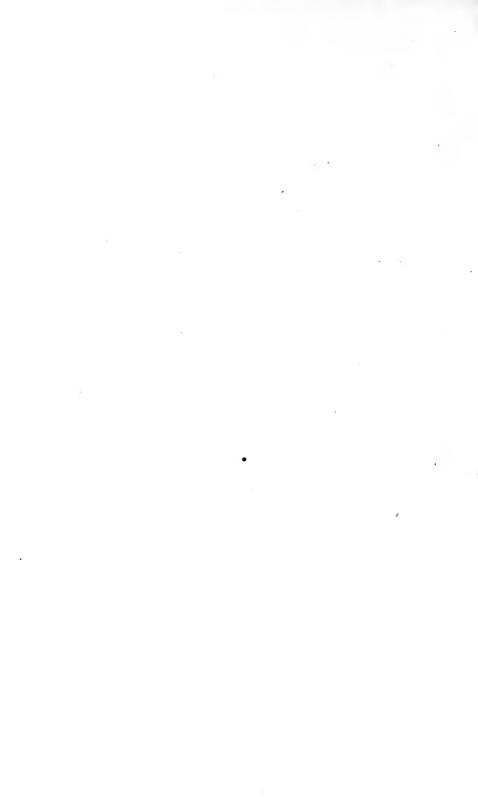
Comment: It is probable that the retailer made a mistake in giving the source of his insect powder, since another sample furnished by Schieffelin & Co. (No. 279) contained no lead chromate.

Serial No. 319.

* * We presume that this analysis is approximately correct, and that therefore there is foreign matter in the insect powder amounting to a little over one-half of 1 per cent. This additional matter is undoubtedly added, not for the purpose of adul-

teration, but for the purpose of brightening the color of the goods. * * * Until recent years it was the custom to sell only colored insect powder, which, of course, required the addition of a considerable percentage of coloring matter. Some years ago, however, we and some others decided that we would endeavor to educate the trade to accept only a strictly pure insect powder, uncolored, and we now supply the colored only, we believe, to one customer, who insists on getting the same. * * * The particular lot of which you sent us the test, however, is not of our own powdering. Last year there was a sudden and very large advance in the price of insect flowers, and we therefore bought in this country the insect powder of a brand that has been held to be one of the most reliable and has had an established reputation for over fifty years. The exceedingly small amount of foreign matter added, amounting to a little over 0.5 per cent, proves conclusively that the object was not to adulterate the powder.

SMITH, KLINE & FRENCH CO.



II. A COMPILATION OF ANALYSES OF INSECTICIDES AND FUNGICIDES.

INTRODUCTION.

In a recent publication b of the Bureau of Chemistry analyses of all the insecticides and fungicides that could be obtained on the American market were given, together with such remarks concerning their effectiveness as the results seemed to justify. It was originally intended that all of the analyses which had previously been made by the various experiment stations should be included in that bulletin. By reason, however, of unavoidable delays, two years passed between the time of collecting the samples and the time of completing the analyses. Since results of this kind, to be of the most value, should be published at once, it was thought desirable to reserve the compilation of results obtained in the various States for a future publication. This article-therefore includes a compilation of analyses of insecticides and fungicides made by the various experiment stations, together with a few analyses made by the Bureau of Chemistry since the publication of Bulletin 68.

The insecticides are arranged in groups, as in the previous bulletin, with the addition of two or three classes not taken up in that publication, and the omission of a few classes not examined by the experiment stations. They will be considered in the following order: (1) Paris greens; (2) London purples; (3) insecticides and fungicides, other than Paris greens and London purples, that contain arsenic, copper, or both; (4) soaps; (5) lyes; (6) hellebores; (7) mixtures containing borax: (8) tobacco extracts; (9) formalins; (10) petroleums; (11) miscellaneous solid insecticides and fungicides, and (12) miscellaneous liquid insecticides and fungicides.

SOURCES OF ANALYSES AND DESCRIPTION OF SAMPLES.

Table I, on the following pages, is introduced to show the publications from which the analyses were taken and, when possible, the names of the manufacturers of the various compounds.

 $[^]a\,\rm No$ responsibility is assumed by the Department for the correctness of compiled analyses, not made within the Department itself.

 $[^]b$ U. S. Dept. Agr., Bureau of Chemistry Bul. 68, The Chemical Composition of Insecticides and Fungicides.

Table I.—Source of analysis and description of samples.

Sample No.	Name.	Source.	Manufacturer of sample.
- 01	Paris tireen	Alabama Experiment Station, Bulletin No.58 (1894)	
100	CD:	op:	
- 10	(d)	000	
: := :	9	96	Eimer & Amend, New York City.
- 0		Cultiformity December of Charles December (1991)	
co	do	do	
. 0		(C)	
=		ор	
21	-	op.	
	(10)	California Experiment Station Report (1895), p. 137	
- 10	District Spiriting Collection	California Decominant Continue Dellocks N. 50	
: :=	American Concentrated Lye	do	
1	Double Concentrated 98 percent Constite Sode	9	Groon Rark Alkali Works Co. Lanoashira England
- 30	Refined Pourl Ashes	O	Do
13	Commercial Potash	op.	Langley & Michaels, San Francisco, Cal.
		op	
_	Paris Green	California Experiment Station, Bulletin No. 126	In New York City.
01		op	
00		op	
-		op:	
	•	ор	
-		op*****	
-		op	
		· · · · · · · · · · · · · · · · · · ·	
n s		(b)	
> -		(10)	
5 84	Concentrated Lye (Potash)	I daho Agricultural Experiment Station. Bulletin No. 19	B. T. Babbitt.
		(1893).	
	American Lye	(a)	American Lye Co., Philadelphia.
	Paris Green (average of 7)	do	
	Sulphur	db	
3 !			
	Taris oredi	dano agricultura experiment station, billicum No. 25	
3	op	(1901).	
23		0)	
0	0	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	
=	ор	00	
Ęį	(10)		

op o	00 00	
	- do - do	
	οpοp.	
	do.	
, 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	op.	
	Not published, but kindly forwarded by Illinois Agri-	A. B. Ansbacher & Co., New York City.
ion brand, new process)	Paris (freen (Lion brand, new process)do	J. A. Blanchard, New York City.
Purigrene	do.	
dodo	op.	H. K. Mulford Co., Chicago, Ill.
op	op	
op	00 do	Fuller & Fuller, Chicago, III.
op	op.	
Formalin (average of 5)	Indiana Experiment Station Report (1899)	Honore Holl f. Co
op.	op.	
do Paris Green (average of 10)	do Louisiana Experiment Station Bulletin No. 1 second	Eimer & Amend, New York City.
Paris Green		Soudheim, Alsberg & Co., New York City.
Paris Green (average of 2)	series (1893), pp. 778–782. Louisiana Experiment Station, Bulletin No. 39, second	
Paris Green (average of 8)	series (1895), pp. 1431–1434. Louisiana Experiment Station. Bulletin No. 45. second	
Pine Tur Inscetteide	series (1896), pp. 80-84. Louisiana Experiment Station, Bulletin No. 48, second	
Paris Green (average of 39)	series (1897). Louisiana Experiment Station, Bulletin No. 49, second	
	series (1897), pp. 198-204. Louisiana Experiment Station, Bulletin No. 54, second	
Paris Green (average of 36)	series (1898). Louisiana Experiment Station, Bulletin No. 58, second	
Paris Green (average of 149)	series (1899). Louisiana Experiment Station, Bulletin No. 63, second	
Paris Green (average of 79)	series (1900).	
	38 (1805).	
	(h)	

Table L.—Source of analysis and description of samples—Continued.

Partic freen (Lien brand, new process) Harch Experiment Station, Massachusetts, Bulletin No. Do. Do.	No.	Same.	Solre.	Mannincurrer of sumple,
100 100	Ś		. Hatch Experiment Station, Massachusetts, Bulletin No. 38 (1896).	Jumes Blanchard, New York City.
Pink Absended		do	90	De.
Fink Arsenaid Hitch Experiment Station, Massachusetts, Builletin No. Arther Color and Chemical Works, New York City, Arther Color and Chemical Works, New York City, Arther Chemical Co., Leoninster, N. Y. Bunderth Chemical Co., Leoninster, N. J. Daniorth Chemical Co., Leoninster, N. J. Arther Chemical Co., Leoninster, Chemical Chem			90	30.
DD DD			Hatch Experiment Station, Massachnsetts, Bulletin No. 74 (1901).	Adler Color and Chemical Works, New York City.
White Arsemeld Do		Green Arsenoid	op.	Do.
Laured Green Chemical Co. Leconimater, N. Y. Paris Green Chemical Co. Leconimater, N. Y. Paris Green Chemical Co. Leconimater, N. Y. Paris Green Chemical Co. Leconimater, N. Y. B. Alusharcher, New York City, Chemical Co. Leconimater, N. Chemical Co. Leconimater, N. Chemical Co. Leconimater, N. Chemical Co. Chemical Chemical Co. Chemical C		White Arsenoid	op	Do.
Part	2!	Lattred Green		Nichols Chemical Co., Syracuse, N. Y.
Triple T	3	Bug Deuth	(10	Danjorth Chemical Co., Leominster, Mass.
Ablaces Abla	3. 3	Paris (ireel)		L. Pfeiller, New York City.
Hampfel Phint and Chemical Co. Hampfel Phint and Chemical Chem			do	Alshers & Pfeiffer New York City
Paris Green (Lion brand) do do do do do do do		,	· · · · · · · · · · · · · · · · · · ·	Hamilden Paint and Chemical Co. Springfield Mass
Paris Green (Linn brand) do	3	:		
Applied to the composite of 2 Applied to the composite of 2 Applied to the composite of 3 Applied to the composite of		Paris Green (Lion brand)	9)	J. A. Blanchard, New York City.
Puris Green do do do do do do do d		ob	olb.	Do,
Description Composition		Paris Green.	ор.	A. B. Ansbacher, New York City.
do	٠.	do	do	Leggett & Bro., New York City.
Paris Green (Lion band, new process)	103	ob	• • • • • • • • • • • • • • • • • • •	E. & F. King Company, Boston, Mass.
Paris Green (Lion brand, new process)	101	ор		C. T. Reynolds & Co., New York City.
Harch Green Composite of 3 Harch Experiment Station, Massachusetts, Bulletin No. Paris Green (composite of 2) Ado Baris Green Brof. De Graffs Bug Destrayer Ado Prof. De Graffs Bug Destrayer Ado Prof. De Graffs Bug Destrayer Ado Prof. De Graffs Green Prof. De Graffs Bug Destrayer Ado Paris Green Paris Green	105	Paris Green (Lion brand, new process)		J. A. Blanchard, New York City.
Paris Green (composite of 3)	-			
Paris Green (composite of 3)	-	:	Hotely Describered Contion Mountainston Dellectin No.	
Paris Green (composite of 3)	-		SI (1919)	
Paris Green do do Paris Green do do Paris Green do do do do do do do d	100	Paris Green (composite of 3)	(10.02):	C. T. Revnolds & Co., New York City.
Paris Green (composite of 2) do do do do do do do d	110	Paris Green	00)	I. Pfeiffer, New York City.
Paris Green	111	Paris Green (composite of 2)	olb	J. A. Blanchard, New York City.
do do do do do do do do	115	Paris Green	op	C. N. Childs & Co., New York City.
do do do do do do do Tobneco Liquor Massachusetts State Experiment Station Report (1890), pp. 275-276. do do do do do do do d	113		ojp	Leggett & Bro., New York City.
Tobacco Liquor Aussichmetts State Experiment Station Report (1899), Ph. 275-276. Ph. 275		op	op	A. B. Ansbacher & Co., New York City.
Tobacco Liquor		ор	(10)	1)0.
do thris Green do the chart of the chart's Bug Destroyer Froi. In c chart's Bug Destroyer day do do do do do the chart's Bug Destroyer do		Tobacco Liquor	 Messachusetts State Experiment Station Report (1890), pp. 275-276. 	
Paris Green Sulplantine Sulplantine Pouth to lase Bugs Prof. De Graff's Bug Destroyer And do	11.7	ob	op	
Sulphutine Death to Rose Bigs Prof. De Graff's Big Destroyer This Green	×	Paris Green		
Sulpuatine Beath to Beath Sang Destroyer Prof. De Graff's Bug Destroyer Paris Green		ф	op.	
Prof. De Graff's Bug Destroyer. Paris Green	-	Sulphatine	36	
Paris Green	3	Death to Rose Dugs	do	
op.	12	Paris Green	Massochnsetts State Experiment Station Report (1892).	
do .			p. 336.	
	124	. 00	do	

	Leggett & Co., New York City. A. J. Peck, Merddian, Miss. Mansfield Deng Co., Memphis, Tenn. I. L. Lyons, New Orleans, La. John Lareas & Co., New York City. Aeme Color Works, New York City. A. B. Anshaderre & Co., New York City. F. W. Devoe & Co., New York City. C. T. Reynolds & Co., New York City. Do.	Hemingway's London Purple Co., London, England. West & Jerenry, Boston, Mass. Leggett & Bros., New York City.	Wetherell & Co. John Lucas & Co., New York City.	Cawley, Clark & Co., Newark, N. J. C., T. Reynolds & Co., New York Giy. Chwley, Clark & Co., New York Giy. Robert Shownaker, N. J. John Lueus & Co., New York Giy. J. A. Blanchard, New York Giy. J. A. Blanchard, New York City. Leggelt & Brost, New York City. Richardson formula. A. Pfeiffer. Leggett & Brost, New York City. Leggett & Brost, New York City.
11111111	1, 324, 1, 3	Organistic for the control of the co	do do Secon Several Station, Bulletin No. 43 Pars Green Several Station, Bulletin No. 43 New Jersey Experiment Station Report (1897), pp. 397- do do do do	Cawley, Clark & Co., Newark, N. Cawley, Clark & Co., New York Cawley, Clark & Co., New York Carl Required & Co., New York Cawley, Clark & Carl & Co., New York Cawley, Clark & Carl & Cawley Cawley, Clark & Cawley Cawley, Cawley Cawley, Cawley Cawley Cawley Cawley Cawley Cawley Cawley Cawley Cawley Cawley
25				102 400

Table 1.—Source of analysis and description of samples—Continued.

C. T. Reynolds & Co. Do. Do. Henj. Hammond, Fishkill on the Hudson, N. Y. Acme Color Works.		C. M. Clark & Co. C. M. Clark & Co. Do. Hampden Paint Co. North Bremann & Co. Do. Pred L. Lavanburg. Do. C. Cavenburg. Do. C. Laverack. Leggett & Bros. Do.	New York Enamed Paint Co. 1 Do. 1 Do. 1 Pfelffer. 2 Do. C. T. Reynolds & Co. Bovoe, Reynolds & Co. Bovoe, Reynolds & Co. Solomon & Solomon & Co. Stanley, Jordan & Co. Stanley, Jordan & Co. Stanley, Jordan & Co. Stanley, Jordan & Co. Stanley forman Sons. Briglish Compound Co. Nichols Chemical Co. Do.
40 40 40 40 60 60 60 60 60 60 60 60 60 60 60 60 60	(1901). (1901)	\$ 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
6 do		2	Company of the control of the contro

Table 1.—Source of analysis and description of samples—Continued.

	Name.	YORKG:	Manufacturer of sample.
	London Purple	New York State Experiment Station Bulletin No. 201 (1901).	Hemingway's London Purple Co.
-	Parngrenc Paris Green, Bordeaux Mixture Discognos	: :3	F. L. Lavanburg. Leggelf, & Bros. Coweker Chemical Co. Roston, Mass.
400.00		op	Swift & Co., Boston, Mass. Laubach & Boyd, Wooster, Ohio.
-			
	Crude Petroleum	op op	
	(1)	00	
_	op		
-	verage of 18)	Oregon Experiment Station, Bulletin No. 49 (1898)	
	Par 'oidiny	do	
	Antinomnin	(P)	
-	Paris Green	South Carolina Experiment Station Report (1888), p. 142	
-	London Purple	do	
	Paris Green	Not published; kindly lurinshed by South Carolina Ex- periment Station.	
-	do	Vermont Experiment Station Report (1888)	C. T. Reynolds & Co., New York City.
_	London Purple		Hemingway's London Purple Co., London, England,
_	Paris Purple	000	A. Porrher, Paris, France.
	Peroxid of Silicates		Peroxid of Sincare Co., New York City.
_	Marie Change	Weekington Perceptores Chatter Propost 1800 as 45 50	naminona s, Fishkin on the Hudson, iv. 1.
	Arsenate of Lead	Not published: data furnished by H. H. Harrington, Texas Agricultural Synchronic Station	F. L. Lavanburg, New York City.
	do.		De.
-	op.	9	Merrimae Chemical Co., Boston, Mass.
	ор.	ор.	Do,
_	op	(I)	Patton-Worshum Drug Co., Dallas, Tex.
_	Paris Green	(d)	Adler Color and Chemical Co., New York City.
-	0.0	0)	Do.
-	op	(Op)	
_	op		
	ojp	(d)	F. L. Lavanburg, New York City.
	London Purple		
	Purale Poison	00	Mover Bros Drug Co. St. Louis Mo.
-	Bollene Insecticide	0)	Bollene Manufacturing Co., Galveston, Tex.
		00	
-	Shig Shot.	do	Benj. Hammond, Fishkill on the Hudson, N. Y.
200	Transmitter		Mol our & Grouns Coldwoll They

			A	MAI	Toro
Eimer & Amend, New York City.	Commercial Lye Co., New York City. E. Schering, Berlin, Germany.	Eimer & Amend, New York City.	Spotteholz & Ameshot, Amsterdam, Holland.	Fly Preventativedododo	Do. J. W. Barwell, Waukegan, III. Baker, Adamson & Co., Easton, Pa.
231 & W Potassium Cyanide Unpublished data of Bureau of Chemistry U. S. Depart- Eimer & Amend, New York City. ment of Agriculture.	ye do do do Gommercial Lye (50, New York City, do do do Gommercial Lye (50, New York City, E. Schering, Berlin, Germany,	and Wood Preserver do	White Arsenie do do do Spotteholz & Ameshot, Amsterdam, Holland	op	645 1 & W do Do. Do
Potassium Cyanide	Royal Concentrated Lye Formalin	281&W Copper Carbonate	1421 & White Arsenie 1431 & W	Fly PreventativeBollene Insectieide	do Karbo-Kresolate Potassiun eyanid (99 per cent)
23 I & W	241 & W 261 & W	281& W 631& W	142 I & W 143 I & W	202 l & W 1 644 l & W 1	6451 & W 9161 & W 11701 & W

PARIS GREENS.

The analyses of a large number of samples of Paris Green are given in Table II, together with notes that aid in explaining the methods of analysis used. Whenever the method of analysis was given in the original publication it is quoted in full, not only in this table, but throughout the bulletin, in order that the reader may judge for himself of its accuracy. In a number of cases in which only one or two constituents were determined in samples of Paris greens the average of all the results is given, mentioning, however, the number that fall below the generally accepted standard for such goods.

METHODS.

The method of determining arsenious oxid in samples Nos. 1 to 6, inclusive, was as follows:

Weigh 1 gram of the material into a beaker, add 30 cc of strong hydrochloric acid and digest at a temperature below the boiling point, adding small quantities of KClO₃ at frequent intervals. Continue heating until the odor of chlorin has disappeared, dilute with water, add ammonia in slight excess, cool, add magnesia mixture drop by drop, stirring vigorously, and let stand twelve hours. Filter, wash with ammonia water, and dry. Detach precipitate from filter and ignite filter, using ammonium nitrate. Transfer the precipitate to a porcelain crucible and heat for a while on an iron plate and finally in the direct flame. Add filter ash and weigh as magnesium pyro-arsenate.

In samples Nos. 37 to 55, inclusive (19 samples), the following method was used for determining soluble arsenious oxid:

Allow 0.5 gram of the green to stand in suspension in 50 cc of distilled water in a stoppered cylinder for one week, with occasional shaking, and titrate an aliquot filtered portion of this solution for arsenious oxid.

The method for determining total arsenious oxid in samples Nos. 69 to 80, inclusive, was practically the same as that used for samples Nos. 1 to 6, inclusive.

In determining total arsenious oxid in samples Nos. 198 to 264, inclusive (62 samples), the method given in Bulletin 68, Bureau of Chemistry, U. S. Department of Agriculture, for the total arsenious oxid in Paris Green was used. In determining copper the electrolytic method was used. In determining soluble arsenious oxid 1 gram was suspended in 1,000 cc of water and allowed to stand over twenty-four hours in a stoppered flask, with occasional shaking. The arsenious oxid going into solution was titrated with iodin solution.

The soluble arsenious oxid in sample No. 285 was determined by extracting 1 gram of Paris Green with 400 ec of water for nine days, with occasional shaking, and titrating the arsenious oxid in an aliquot portion of the resulting solution with iodin.

Table II.—Composition of Paris greens.

Remarks.	This sample contained neither copper nor arsenic, but consisted of a mixture of prussian blue, chrome yellow, and either clay	OF CHAIK.	This sample also contained a trace of potassium oxid, 0.58 per cent zine oxid, and 0.75 per cent of prussian blue. The manufacturers of this Paris green claimed 64 per centarsculings and 90 new cent of contact oxid and 90 new cent of contact oxid.		In sample No. 34 the lighter for accentance oxid is an average of 7 lawris greens, one of which contained below 50 per cent arse, nious oxid, having only 39.61 per cent. The figure for copper oxid is an average of 2 laris greens.	
Soluble oxid in arsenious oxid in oxid (As ₂ O ₃), copper.	Per cent.					
Soluble arsenious oxid (As ₂ O ₃).	Per cent.		23.60 29.40 24.60	6, 70 8, 10 9, 10	8. 2. 2. 3. 44. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5. 5.	######################################
Caleinm oxid (CaO).	Per cent.		0.10	2 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		
Sodium oxid (Na ₂ O).	Per cent.		0.77 Present.	1.51 1.01 38.		
Silien (SiO ₂).	Per cent.		0.10	.13		
Sulphur trioxid (SO ₃).	Per cent.		0.23 Present.	1.30		
Acetie acid.	Per cent. Per cent.	11.45 11.15 8.95 30.15	11.28 0.23 Present.	3.93		
Total copper oxid (CuO).	Per cent.	52.5	31.25 32.60 26.70	21.81.81.81 88.82.81	23. 25. 25. 25. 25. 25. 25. 25. 25. 25. 25	58.58.58.58.58.58.58.58.58.58.58.58.58.5
Total arsenious oxid (As ₂ O ₃).	Per eeat. 55, 42 55, 42 55, 71 53, 13 57, 38 54, 15	2.5.5.2.3.3.2.0.2.3.3.3.3.3.3.3.3.3.3.3.3.3.3	52.90 55.13	4.8.8.9. 72.8.8.2.2.	55. 69 55. 08 57. 07	8 8 8 8 2 2 2 8 8 8 8 8 8 8 8 8 8 8 8 8
Moisture, arsenious oxid (As ₂ O ₃).	Per cent.	20.1.68.8.2.9.1.9.1.9.1.9.1.9.1.9.1.9.1.9.1.9.1.9	1.31	2.00		
Sam- ple No.	1284595	*62122	គ សភ	<u> </u>	£ 25 25 25 25 25 25 25 25 25 25 25 25 25	84444444

Table II.—Composition of Paris greens—Continued.

		This figure is the average of 10 Paris greens, 2 of which contained less thm 50 per cent arealions oxid, viz. (1), 41.28 per cent, numufactured by G. T. Reynolds & Co., New York City; (2), 49.82 per cent, nanufacturer unknown.	Average of 2 Paris greens, one containing less than 50 per cent arsenious oxid (48.47 per cent, manufactured by Weiland & Meissner). This figure is the average of 8 Paris greens. This figure is the average of 39 Paris greens, 1 containing less than 50 per cent arsenious oxid (48.63 per cent, manufacturer unknown).	Average of 3P Paris greens, 1 contained less than 50 per cent arsenions oxid (49.69 per cent, obtained of Martin & Ducote, Lathyyette Jowa). Average of 3D Puris greens. Average of 19 Puris greens. Average of 3P Puris greens. Average of 3P Puris greens. Average of 3P Puris greens.	arsenious oxid (49.38 per cent, manunacturer unknown). This sample also contained 0.37 per cent of magnesium oxid.
Arsenious oxid in combina- tion with copper.	Per cent.				
Soluble arsenious oxid (As ₂ O ₃).	Per cent.	æ :			14.50
Calcium oxid (CaO).	Per cent.				
Sodium oxid (Na_2O) .	Per cent. Per cent. 21.31				
Silica (SiO ₂).	Per cent.				1.40
Sulphur trioxid (SO ₃).	Per cent. Per cent. Per cent. Per cent. 21.31				7,56 1,40
Acetic acid.	Per cent, Per cent.				
Total copper oxid (CuO).	Per ex 18 18 18 18 18 18 18 18 18 18 18 18 18	9. S			30 30 35 35 35 35 35 35 35 35 35 35 35 35 35
Total arsenious oxid (AsyOg).	74 888888888888888888888888888888888888	52. 55 52. 55 53. 55		7.17 58.59 58.73 58.73 58.73	15.12.85.88 8 = 15.94 8 = 15.94
am- pple Moisture, arsenious No. (AsyO.).	Prent.		*		8.6.5.9.8 8.6.5.9.8
No.	#2282222222222 #25252222222222	38 8	7,5 7,5	853313 n	23818

	ANALISES OF	INDE	CITCIDES	AND FUN	GIOIDES.	0.
This sample also contained 0.56 per cent of magnesium oxid. This sample also contained 1.15 per cent of magnesium oxid and 0.94 per cent of potassium oxid. This sample also contained 1.37 per cent of magnesium oxid and 0.03 per cent of potassium oxid.		This sample is a composite of 3 greens manufactured by the same person.	This sample is a composite of 2 samples manufactured by the same firm.		In reporting samples Nos. 138-141, inclusive, the chemist reported the copper as copper hydric, added the copper hydrate, moisture, and arecnious acid together and subtracted fron 100, reporting the remainder as impurities. The acetic acid was not taken into account.	This figure is an average of 5 Paris greens.
0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0						
					1.09	1.09
15.98 15.67 16.88						
i 38 53						
09		0.01	1. 20.1.20 2.	358388		
5.78						
			8.93	2.00 2.00 1.7 2.00 3.00 3.00 3.00 3.00 3.00 3.00 3.00		
8. 96 7. 88	**************************************	28.08 29.08 29.08	88 88 88 88 8 24 88 88 88 8	18.88.88.88.88 18.89.98.88 18.89.99.88		
63.98 47.52 49.70	88488888888888888888888888888888888888	59,04 61,15 60,38	6.66.66.21 58.66.61 2.85.82	8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	56. 55. 17. 17. 17. 17. 17. 17. 17. 17. 17. 17	59.55
.71		8'8' B	85. 7.8 39. 39. 39. 1.31	11.12 1.15 1.15 1.15 1.15 1.15 1.15 1.15		
£ & & &	58828888888 5882888888	109	11 11 11 11 11 11 11 11 11 11 11 11 11		29 <u> </u>	147

Table II.—Composition of Paris greens—Continued.

Remarks,	The 43.3 per cent, insoluble in hydrochloric acid, consisted of 30.6 per cent. silica, 5 per cent oxids of iron, and aluminum and other constituents. This green was evidently composed of about equal parts of Paris Green and carthy matter, such as elay or infusorial earth.								This figure is the average of 5 Paris greens.		These figures are the averages of 24 Paris greens.								7.											
Arsenious oxid in combina- tion with copper.			:			:	:			:	:	:	:									:	:							_
Soluble arsenious oxid (As ₂ O ₃).	Per cent. Per cent.									1.83		1.35	1.10		3.5	1.10	15.69	7.7	7	1, 10	1.23	86.	98.	22.	6.6	8.5	3	S. 32	98.	000
Calcium oxid (CaO).	Per cent. Per cent.		:			:	:				:		:	:								:		:	:	:				-
Sodium oxid (Na ₂ O).						:						:	:	:			:						:	:	:					
Silica (SiO ₂).	Per cent.		:											:									:	:	:	:				
Sulphur trioxid (SO ₃).	cent. Per cent. Per cent.									Ξ.				:											:					
Acetic acid.	Per cent.					:						:	:					:					:							
Total copper oxid (CuO).	Per		:			:				33.65	29.92	30.30	50 S	30.3	8 8 8 8 8	29, 98	98.9	25, 55	30.53	28.76	30, 16	30,39	30,16	30.39	23.45	30.70	95.10	30.70	30, 39	100 000
Total Total Moisture, arsenious oxid (As ₂ O ₃).	P.	56.13 56.19		\$ -	56.03	63.11	66.43	59.33	54.73	56,45	56.48	56.56	26.30	56.55	5.5. 56.8.5	56.75	59.20	26. 60	57.17	57, 11	56, 62	57.11	56.50	36.93	00.70	25.3	9	57.05	56, 75	27. 12
Moisture.	Per cent.		:									:	:					:						:						_
No.	159	353	163	2 2	991	167	2 2	170	121	2	195	8	200	900	200	503	501	500	202	503	503	510	211	775	7 5	7 5	216	217	218	010

25.27 55.24 56.24 66.24	56.38	52,53	25.10	53.46		55.56	25.00	57.42	55.95	56.97	26.30	51.46	-	F	_	_	because the present New York	_	_	_	57.13	56.97	56.27	56.63	96. 85	56.27	57.57	56.74	95.11	52.23	00.75	01.76	955.45	17 'GG	57.72	oc. oc	•	The state of the s	The ngure 24,53 is the average of 18 Paris greens, 5 of which are	Defow 50 per cent arsenious oxid (viz, 46.50 per cent, 57.30 per cent, 45.51 per cent, 45.08 per cent, 40.98 per cent, 46.16 per cent,	34.10 per cent, 46.60 per cent; manufacturers unknown). The	figure 29.47 is the average of 12 Paris greens.			
		138		1.53	_					_	1.10													1.72			88.	_				_		_		S.	:	:	:				88 6		40 II.42 ITREE.
		_				_	-							_	_		_	_	_	_	_	_	_		_		-	_		,					_	•	:	:	:						
																																				:		:							
		:																									:		:	:	:	:	:	:	:	:		:	:						
		:	:					-		-							-								:				:	:	1	:	-	-	:	:	:	:	:				:	Tracoo	Truce
		:																									:	:		-	:	:	:	-		:	:	:	:						:
		:																									:	:	:	:	:	:	:	:	:	1	2:		:				:		:
																													:		:	:	:	:		:	8 i		:					=	11.42
30,0		8.8				99.66			30.08			27. 47		26. 53														30.29											29, 17				68 08	90.07	
57.66	56.93	57.11	56.01	57. 49	57 11	57, 35	57 79	57. 42	57 85	58.76	30	9	62.87	65.69	57.91	57.91	30	58.03	57.11	56, 13	57.42	57.42	57.11	58.09	58.46	57.35	57.72	58.09	60.72	57.29	57.60	Z.	50,00	57.60	57. 75	50.00	55.55 50.55	98. 99	5 1 . 4 3			00	98.99	20.00	50.76
		:		:						:	:						-	-						:	:	:	:::::::::::::::::::::::::::::::::::::::	:	:	:					:		:		:	-			2	1 10	1.10
* *																																													

Table II.—Composition of Paris greens—Continued.

Remarks.	Analysis made in accordance with Texas law regulating sule of Paris greens and other insecticides and is only good until	May 1, 1965. Analysis made under Texas law; good only until May 1, 1902. Do. Do.
Arsenious oxid in combina- tion with copper.	Per cent.	25.25.21.25.25.25.25.25.25.25.25.25.25.25.25.25.
Soluble Arsenious oxid in coxid tion with (As ₂ O ₃), copper.	Per cent. 0.95	2.7.3.5. 2.7.3.9.
Sodium Calcium Socid Oxid (NagO). (CaO).	Per cent.	
Sodium oxid (Na ₂ O).	Per cent.	39,00 29,17 33,20 29,05
Silica (SiO ₂).	Per cent.	
Sulphur trioxid (SO_3) .	Per cent.	
Acetie	Per cent.	30, 00 29, 74 31, 20 29, 65
Total copper oxid (CuO).	Per cent. 29, 77	30,00 31,23 29,63 65,63
Sam- ple Moisture, arsettious No. (Asj.O ₃).	Per cent. 55, 90	25 25 25 25 25 25 26 25 25 26 25 25 26 25 25 26 2
Moisture,	Per cent.	
Sam- ple No.	196	86.69

COMMENT ON ANALYSES.

A glance at Table II at once makes it apparent that only 1 sample out of more than 650 examined contains neither copper, acetic acid, nor arsenious oxid—this is sample No. 7. One other sample, No. 159, consists of about equal parts Paris Green and some insoluble material that appears to be clay or infusorial earth. Out of the total number examined 31 contained less than 50 per cent arsenious oxid. It will be noted, however, that a number of these samples said to contain less than 50 per cent arsenious oxid were analyzed by a method (the magnesium pyro-arsenate method) that has since been shown to give results about 1 per cent low. If a more accurate method of analysis had been used, the number falling below 50 per cent arsenious oxid would very likely be materially diminished. It also appears that the Western States receive more of those samples that contain small percentages of arsenious oxid than do the Northern States.

The State of Louisiana, which previous to 1899 occasionally received samples containing small percentages of arsenious oxid, does not seem to have had any upon the market during 1899 and 1900. Out of 96 samples of Paris Green examined in New York State during the years 1899 to 1901, inclusive, none was found that contained less than 50 per cent arsenious oxid. In both California and Oregon, until a very recent date, quite a large amount of the Paris Green on the market contained small percentages of arsenious oxid, although it is said that this condition has been very much improved during the last two years.

Concerning soluble arsenious oxid in Paris greens, it appears that those upon the market in California until recently contained very large quantities, those in Idaho were very fair, only 2 out of 19 containing more than 4 per cent, those in Illinois, Montana, and Texas contained very small percentages, and those in New York the same, only 1 out of 90 containing more than 4 per cent. This one was sample No. 204, containing 15.69 per cent of free arsenious oxid.

It must be borne in mind, however, in looking over the figures for free arsenious oxid that the results in one State can not be compared with those in another, since the methods of analysis were very likely different in every case and would lead to entirely different results. It may be of interest to those who have not read the previous bulletin of the Bureau of Chemistry on this subject, to mention that out of the 45 samples of Paris Green, from all parts of the United States, which were examined 71 per cent contained more than 4 per cent of soluble arsenious oxid and 16 per cent contained more than 6 per cent of arsenious oxid when a ten-day extraction of 1 gram with 1,000 cc of water was made. In view of the facts brought out in that bulletin, a 6 per cent, rather than a 4 per cent, limit for soluble arsenious oxid was advocated when the above method of analysis was employed.

From the figures given above, together with information received from the various States, it appears that upon the whole the purity of the Paris greens upon the American market is steadily and rapidly upon the increase.

LONDON, ENGLISH, AND PARIS PURPLES.

The analyses of 10 samples of London Purple, 1 sample of English Purple, and 3 samples of Paris Purple are given in the following table All of these are products obtained by boiling a waste product from the analine-dve industry with lime. Since it was not known at the time these analyses were made that the arsenic was present, both as arsenious and arsenic acids, the total arsenic was determined and reported as arsenious oxid, only the soluble arsenious oxid being determined, instead of both soluble arsenic and arsenious oxids.

Table III.—Composition of London purples and closely allied mixtures.

Sam- ple No.	Name.	Moisture.	Arsenious oxid (As ₂ O ₃).	Calcium oxid (CaO).	Ferric oxid and alumina $(Fe_2O_3$ and $Al_2O_3)$.	Sulphur trioxid (SO ₃).	Dye.	Soluble arsenious oxid (As ₂ O ₃).
	London Purple		37.18		Per eent.			Per eent. 0.54 2.58
175	do dodo	. 3.27	41.44	24.32	3.37		27. 29	
268 284	dodododo	3.99	32.32 42.60					12.21
302	dodododo				1.31			
178 1 179 1	English Purple Paris Purple		36.75 47.05					5. 36 11. 86
	do		34. 10 33. 72	4.55	4.76			

a A portion digested 40 hours with water at room temperature gave 8.11 per cent arsenious oxid and a A portion digested 40 hours with water at room temperature gave 8.11 per cent arsenious oxid and 2.85 per cent soluble calcium oxid. A portion digested with hot water until all the soluble arsenious oxid had dissolved gave 16.45 per cent soluble arsenious oxid and 3.26 per cent soluble calcium oxid.
b Also contains 1.65 per cent nitrogen.
c This analysis was made in accordance with the Texas law regulating the sale of Paris Green and other insecticides and is good only until May 1, 1903.
d Made in accordance with Texas law, but good only until May 1, 1902.
c Also contained 2.8 per cent nitrogen and 0.57 per cent potassium oxid.

It will at once be noted that the variation in the amount of arsenic in different samples of London Purple and Paris Purple is very large. In the case of 11 samples examined more recently in this Bureau (published in Bulletin 68) this variation in the arsenic content is not nearly so great. It would, therefore, appear that the amount of this constituent in such goods is becoming more constant in the more recently manufactured samples.

The figures for soluble arsenious oxid can not be compared with those obtained more recently in this laboratory, since the soluble arsenic oxid was not determined in the earlier analyses, and it is only when both figures are at hand that any conclusions of a positive nature can be drawn.

INSECTICIDES AND FUNGICIDES, OTHER THAN PARIS GREEN AND LONDON PURPLE, THAT CONTAIN ARSENIC, COPPER, OR BOTH, AS THEIR ACTIVE CONSTITUENTS.

In Table IV is given a compilation of analyses of Green Arsenoid, Green Arsenite, Paragrene, Laurel Green, and White Arsenic, together with two analyses of White Arsenic made in this laboratory.

Table IV.—Composition of Green Arsenoid, Green Arsenite, Paragrene, Laurel Green, and White Arsenic.

Moisture,	Total arsenious oxid (As ₂ O ₂).	Copper oxid (CuO).	Calcinm oxid (CaO).	Sulphur trioxid (SO ₃).	Soluble arsenious oxid (As ₂ O ₅).
Per cent.	Per cent.	Per cent.	Per cent.	Per cent.	Per cent.
2.77	61.43	28, 83			7.82
1.44	50.77	31.90			
	60, 63	29. 29		1.55	3.53
	53.71	29.43		3.70	8.33
1.91	58, 82	30.76			2.9
	54.10	31.59		3.70	2.36
6.20	40.60	23, 46			23.08
	19.31		13, 31		
	55, 57	27, 68		1.34	3, 21
	43.34	18.08			
	52, 30	21.64			
8. 15	36, 11	17.87	14.20		1.4
7.64	7.34	13, 50	26, 31		
	7.75	12.52			
ecial)	42, 69	4, 50	35, 39		
	3, 83	11.50			
		12.68			. 00
		12.05			
	98, 00				
	99 99				
•		99. 86 99. 99	99.86	99.86	99.86

a The following constituents were also present: Silica, 0.40 per cent; organic matter, 6.67 per cent. b In sample No. 30 the following constituents were also present: Acetic acid, 6.72 per cent; calcium sulphate, 19.31 per cent; sodium sulphate, 2.26 per cent; sodium chlorid, 0.25 per cent; ferric oxid, 6.20 per cent.

C Large amounts of calcium carbonate and hydrate were also present.

The figures obtained by different analysts for Green Arsenoid, Paragrene, and Laurel Green show that the composition of these substances is very variable, especially as to the content of arsenious oxid, which varies in the first case from 50.77 per cent to 61.43 per cent, in the second from 19.31 per cent to 55.57 per cent, and in the third from 3.83 per cent to 42.69 per cent. The figures obtained for soluble arsenious oxid, here as elsewhere in this compilation, can not be compared, since the methods of extraction were very likely different.

The two samples of White Arsenic analyzed in this laboratory were received from Boise City, Idaho, by the Division of Entomology. They are both evidently of exceptional purity and will give good results when used for preparing Calcium Arsenite, the purpose for which they were purchased.

d This analysis was made in accordance with the Texas law regulating the sale of Paris Green and other insecticides and is good only until May 1, 1902.

The method of analysis used is that described for total arsenious oxid in Paris Green in Bulletin 68 of this Bureau.

WHITE ARSENOIDS.

Table V.—Composition of White Arsenoids,

Constituents.	Sample No. 27.	No. 91.
	Per cent.	Per cent.
Barium oxid	42.39	41.43
Barium to join with chlorin Lead oxid	8,60	6. 10
Lead oxid	1,55	. 90
Carbon dioxid	10.15	8, 9:
Chlorin	4.45	3.19
Silica	. 20	
Arsenious oxid	27, 64	31.96
Moisture	4.00	2.3
Soluble arsenious oxid	27, 64	

PARIS GREEN AND BORDEAUX MIXTURE.

The composition of this sample, No. 270, was as follows:

Pe	er cent.
Total arsenious oxid	15.49
Copper oxid	16.02
Soluble arsenious oxid.	

SLUG SHOTS.

Table VI.—Composition of Slug Shots.

Constituents. Sample No. 181. Sample No. 2	le Sampl 90. No. 307
Per cent. Per ce	
	60
hur trioxid 43	. 25 28. . 02 41.
ic oxid and alumina	79 83
luble	38
tile organic matter (by difference).	

The sodium chlorid in sample No. 290 was calculated from a content of 1.72 per cent of chlorin. The volatile organic matter in this sample consisted principally of dead oil.

In sample No. 307 small amounts of sulphur, copper sulphate, oxid of iron, carbolic acid, and tobacco, with less than 1 per cent of arsenic, were also present.

Slug Shot is evidently a mixture of fairly constant composition containing in all cases small amounts of arsenious oxid and copper oxid as active poisonous constituents.

PEROXID OF SILICATES.

Table VII.—Composition of peroxid of silicates.

	Constituents.	_	Sample No. 135,	Sample No. 289.
			Per cent.	Per cent.
Moisture			1,65	1.4
Arsenious oxid			. 57	1.4
Copper oxid			. 33	. 2
ulphur trioxid			49.66	52.2
aleinm oxid			41.18	36.4
nsoluble			2, 31	1.9
erric oxid and alumina				1.4
	(by difference)			4.7

This mixture evidently contains arsenious oxid and copper as the active constituents, diluted with a large amount of gypsum. In so far as the arsenious oxid and copper content are concerned, this mixture is of value as an insecticide; but the gypsum is inert in its action upon insects, except that when the spiracles are not well protected they may be closed up with this or any other fine dust.

BLACK DEATH.

Composition of Black Death.

[No. 222.]	
[50, 222.]	Per cent.
Moisture	9.78
Sulphate of lime (gypsum)	45.34
Magnesium oxid	3.98
Ferric oxid and alumina	3.02
Arsenious oxid	
Copper oxid	41
Silica	5.42
Loss on ignition	28.91

ENGLISH BUG COMPOUND.

The composition of this compound, No. 265, is as follows:

Per	cent.
Arsenious oxid	1.46
Copper oxid	. 60

ORIENTAL FERTILIZER AND BUG DESTROYER.

[No. 136.]

Composition of Oriental Fertilizer and Bug Destroyer.

	Per cent.
Water	37. 14
Total solids	12. 86
Arsenic oxid	2.38
Potassium oxid	3.50
Sodium oxid	6.08
Nitrogen	67
Chlorin	3.00
Sulphur trioxid	64

This mixture is evidently sodium and potassium arsenates dissolved in water. In so far as its content of arsenic is concerned, it would be of value in killing insects. It might, however, if sprayed upon trees, cause some damage by scorching, as the arsenic is in a soluble condition. Its fertilizing value is shown by its content of potash and nitrogen.

Another sample of Oriental Fertilizer and Bug Destroyer, No. 150, gave, in 1,000 cc of liquid, 13.12 grams of arsenic and 10.28 grams of potassium oxid.

LEAD ARSENATE AND PINK ARSENOID.

The analyses of samples of Lead Arsenate and Pink Arsenoid, or Lead Arsenite, are given in the following table:

Table VIII.—Composition of Lead Arsena'e and Pink Arsenoid.

Sample No.	Name.	Moisture.	Lead oxid (PbO).	Arsenie oxid (As ₂ O ₅).	Arseni- ous oxid (As_O ₃).	Soluble arsenious oxid (As ₂ O ₃).
a 271	Disparene	Ter cent. 40.40	38.50	13, 28		
b 272 c 292	Lead Arsenatedo	29.00		14, 83	d 21, 20	
e 294	do do do	f 35, 30 f 31, 50 f 37, 40	32.34 44.40 44.98		d 18,00	
e 295	do	32, 92	34. 83 49. 58		d 30, 62	
89	dodo	.35				1.93

a All the lead oxid calculated to Pb₃(AsO₄)_a leaves an excess of 0.05 per cent arsenic oxid, probably combined as sodium arsenate.

It is at once evident that the variation in the composition of samples of Lead Arsenate is considerable, depending to a large extent on the variation in the moisture content and somewhat on the variation in method of manufacture. These differences in composition should be borne in mind in using such compounds for spraying purposes, since some of them would not have the same insect-killing powers as others at the same dilution. Pink Arsenoid, although somewhat variable in composition, is much more constant than is Lead Arsenate.

XX, PURPLE POISON, AND INSECTICIDE.

Table IX.—Composition of XX, Purple Poison, and Insecticide.

Constituents.	XX (No. 187).	Purple Poison (No. 304),4	Insecticide (No. 308),a
Total arsenious oxid		30, 6 8, 5	
Lead oxid (combined)			66, 60
Organic matter (volatile)	32, 23	9.5	9, 50 8, 00

[&]quot;The Purple Poison, sample No. 304, also contains coloring matter. The two analyses, Nos. 304 and 30s, were made in accordance with the Texas law regulating the sale of Paris Green (see page 61) and other insecticides and holds good only until May 1, 1903.

b All the lead oxid calculated to $Pb_3(AsO_4)_2$ leaves an excess of 1.38 per cent arsenic oxid, probably

b All the lead oxid calculated to Pb₃(AsO₄)₂ leaves an excess or 1.05 per cent atsente oxid, probably combined as sodium arsenate.

c This analysis was made in accordance with the Texas law regulating the sale of Paris Green and other insecticides and is good only until May 1, 1903. In addition to constituents named, 2.7 per cen organic matter was found.

d The chemist reported arsenic in the following ways on these 5 samples: (1) White arsenic, (2) total arsenic, (3) white arsenic, (4) white arsenic, (5) arsenious oxid. These terms are usually applied to arsenic when in the "ous" condition and mean so much As₂O₃. If these compounds are arsenates, as claimed, the arsenic should be reported as As₂O₃. In several of the above analyses, however, if the figures as given were calculated to As₂O₅, the sum total of the constituents would be markedly above 100 per cent.

e Analysis made according to Texas law, but good only until May 1, 1902.

Analysis made according to Texas law, but good only until May 1, 1902.

f Includes organic matter also.

g In addition to constituents named 6.85 per cent of organic matter, lead sulphate, etc., was found

None of the three analyses is sufficiently complete to show definitively the chemical composition of the compound examined, although from the data as given the Purple Poison appears to be at least partially composed of London Purple or some closely allied product.

COPPER CARBONATE.

[No. 28 I and W.]

The analysis of this sample in the Bureau of Chemistry showed that it contained 58.24 per cent of total copper oxid. It is basic in character and of somewhat variable composition, so that the purity can not be calculated from the copper-oxid content. It contains, however, a somewhat smaller amount of copper oxid than other samples examined in this laboratory. The method of analysis used was that described in Bulletin 68.

FOSTITE.

Composition of fostite.

[No. 280.]	
Pe	er cent.
Loss at dull-red heat	2.92
Soluble in aqua regia:	
Copper oxid	2.98
Ferric oxid and alumina	1.36
Calcium oxid	2, 31
Magnesium oxid	1.51
Sulphur trioxid	3.28
Silica	. 35
Insoluble in aqua regia	83.77
Soluble in hot water	7.49

This material is said by the analyst to be composed of copper sulphate and finely powdered soapstone in the ratio of approximately 1:9.

SULPHATINE AND DEATH TO ROSE BUGS.

Table X.—Composition of Sulphatine and Death to Rose Bugs.

Constituents.	Sulphatine (No. 120).	Death to Rose Bugs (No. 121).
Moisture Calcium oxid Copper oxid	Per cent. 1, 40 18, 60	Per cent. 2, 95 17, 76
Sulphur trioxid Sulphur. Insoluble	4.73 48.28 1.63	1.05 4.35 34.53 .49
Ash		54. 14

Neither of the above substances has ever been examined in this laboratory, but from the analyses it would appear that sulphur and copper, probably as copper sulphate, are the active constituents. These are diluted with inert materials, probably to mitigate their action upon the foliage.

SOAPS.

Table XI.—Composition of whale-oil soaps.

Sample No.	Moisture.	Potash lye.	Soda lye.
171	Per cent. 27.30 71.58 6.22	Per cent. 10, 55 6, 86 13, 86	Per cent. 7, 54 4, 89 9, 90
174		10.66	7.61

The above analyses do not indicate that both soda and potash are present, but show how much of either would be present calculated from the number of cubic centimeters of normal acid necessary to neutralize a known weight of soap.

LYES.

The compiled analyses of a number of samples of lyes and the results on one sample (No. 24 I & W) examined in this laboratory are given in Table XII.

Table XII.—Composition of lyes.

Sample number.	Name.	Mois- ture.	Sodium hydrate (NaOH).	Potassium carbonate (K ₂ CO ₃).	Sodium ehlorid (NaCl).	Sodium carbo- nate (Na ₂ CO ₃).	Sodium sulphate (Na ₂ SO ₄).	Potassium hydrate. (KOH).	Potas- sium sul- phate. (K ₂ SO ₄)
15	Solid Lye	Perct.	Per cent, 22.7	Per cent.					
16	Concentrated Lye Double Concen-	.5			5. 9		13. 2		
18	trated Caustic Soda Refined Pearl		99.3						
1	Ashes	23.8		69.1					
	Commercial Pot- ash	1.5		16.8	4.9		14.8	52.3	7. !
	Potash Concentrated	4.5		71.4		2.3			14.
33 °	Potash Lye Concentrated Lye	.58 1.87			7.56 16.70	.33	4, 85 4, 94		
	Royal Concen- trated Lye		70, 96			3. 17			

a The following constituents were also determined: 2.1 per cent insoluble, and 0.1 per cent silica.

The method used for the analysis of the Royal Concentrated Lye (No. 24 I & W) is as follows:

The lye is weighed in a weighing bottle, dissolved to a definite volume in carbon-dioxid-free water and aliquot portions taken for analysis. Phenolphthalein is added to one of these portions and N/3 potassium acid sulphate is added at the rate of about a drop per second, with constant stirring until the pink color fades and the solution becomes colorless. The number of cc of N/3 acid used, "n" represents the

bThe following constituents were also determined: 3.6 per cent potassium chlorid, and 2.7 per cent insoluble.

cThe following constituents were also found: 0.06 per cent of alumina and traces of ferric oxid, magnesium oxid, and insoluble.

sodium hydroxid and one-half the sodium carbonate, since the sodium carbonate is changed to sodium bicarbonate (NaHCO₃). A drop of methyl orange is added and the titration continued until the appearance of a pink color. Let this number of cc be represented by "m." Since this last titration represents the sodium bicarbonate present or one-half the sodium carbonate, "2 m" represents all the sodium carbonate, and "n-m" the sodium hydroxid.

The figures in most cases explain themselves, although a few remarks emphasizing certain points might not be out of place. Sample No. 15 is evidently a very poor example of a soda lye, as two-thirds of it is composed of sodium chlorid. No. 16 is a very fair sample of caustic soda, while No. 17 is a most excellent sample of the same goods. No. 32 is not a potash lye, as it claims to be, but a soda lye. Nos. 32, 33, and 24 I & W are all fair samples of commercial soda lye.

HELLEPORES.

Table XIII.—Composition of hellebores.

Sample No.	Ash insol- uble in hy- drochloric acid.	Total ash.	Sample No.	Ash insoluble in hydrochloric acid.	Total ash.
133	Per cent. 2,34	Per cent.	156	Per cent.	Per cent.
134 154	38.12	33. 13 16. 00	157 158		10.6 34.0

Samples Nos. 134, 154, and 158 are evidently badly contaminated with dirt, probably because the roots of the plant were not properly cleaned. Sample No. 155 also appears to have a slightly higher ash content than should be allowed in preparations of this kind.

MIXTURES CONTAINING BORAX.

A single example of this class of compounds, the analysis of which was made in this laboratory, is given below:

HOOKER'S COCKROACH AND WATER BUG EXTERMINATOR.

Composition of Hooker's Cockroach and Water Bug Exterminator.

	Per cent.
Camphor	3.5 - 4.0
Borax (Na ₂ B ₄ O ₇ 10 H ₂ O)	92, 25
Ammonia (NH ₃)	. 27

The remainder is principally organic matter and appears to be tobacco. The ammonia seems to be present as ammonium carbonate.

In making an analysis of this compound the camphor was determined by extracting the powder with ether, allowing it to evaporate at room temperature and weighing at once. While this method leaves

much to be desired, it at least gives approximate results when a large portion of the material is used. The borax was determined by the same method as given in Bulletin 68 for similar compounds, except that the ammonia was all driven off from the solution by boiling before the boric acid was determined.

This mixture would very likely be efficacious in driving away cockroaches and water bugs, but probably it would not kill them.

TOBACCO EXTRACTS.

The analyses of five samples of tobacco extracts are given in Table XIV. In two of these only the most important constituent, nicotin, was determined, while in one of them the determination of this constituent was not made.

Table XIV.—Composition of tobacco extracts.

Constituents.	Sample No. 116.a	Sample No. 117.b	Sample No. 130.	Sample No. 131.	Sample No. 132
	Per cent.	Per cent.	Den soud	Per cent.	Pêr cent
(fairence					201 00111
Moisture		40.890			
Ash		27, 770			
Fotal nitrogen	2.010	1.730			
Nitrogen as nitrates					
Nicotin	2.115	. 530	4.55	4.82	
Ferrie oxid and alumina	229	. 017			
Calcium oxid	3, 069	1.466			4.
Magnesium oxid	2,303	1, 121			
Phosphorus peutoxid	404	. 057			
Sodium oxid	207	.525			
Octobrish oxid		16, 340			
Potassium oxid	0.000				
Insoluble	Trace.				2.

a Specific gravity, 1.3858.

In the first four of these samples the nicotin figure is much too low for a good tobacco extract; especially is this true of the first two.

FORMALINS.

The analysis of three samples of formalin, and the average of the results obtained on five samples, together with one analysis made in this laboratory, are given in the following table:

Table XV.—Composition of formalins.

	Formal	dehyde.
Sample No.	By weight.	By volume
•		Per cent.
5	36.9 37.0	40. 40.
7s	38 9 a 36, 69	41.
6 (I & W)	37.18	

b Specific gravity, 1.3777.

The method used for determining formaldehyde in samples 65 to 68, inclusive, is as follows: \dot{a}

Fifteen cubic centimeters of $N/10~AgNO_3$ are treated with 5 drops of 50 per cent nitric acid in a 50 cc flask. Ten cc of a solution of potassium cyanid (containing 3.1 grams of 96 per cent potassium cyanid in 500 cc of water) are then added, the flask filled to the mark with distilled water, well shaken, and an aliquot portion of the filtrate, say 25 cc, titrated according to the method of Volhart with a N/10 solution of ammonium sulphocyanate for the excess of silver.

Another 15 cc portion of N/10 AgNO₃ is treated with 5 drops of 50 per cent nitric acid and 10 cc of the potassium cyanid solution, to which has been added a known amount of the formalin solution. The whole is made up to 50 cc, and a 25 cc filtrate from it titrated with N/10 ammonium sulphocyanate, the same as before for the excess of silver. The difference between these two results gives the amount of potassium cyanid that has been used by the formaldehyde in terms of N/10 ammonium sulphocyanate, and since each cubic centimeter of this is equal to 3 milligrams of formaldehyde, we can obtain the amount of formaldehyde that has been acted upon by the potassium cyanid.

The method used in the analysis of sample No. 26, examined in this laboratory, is as follows:

About 3 grams of the solution is weighed into a tall Erlenmeyer flask containing 25 to 30 cc of double normal sodium hydroxid. Fifty cc of pure 2.5 to 3 per cent fresh hydrogen peroxid is then gradually added, during a space of three minutes, through a funnel placed in the neck of the flask to prevent spurting. After standing ten minutes the funnel is washed with water and the unused sodium hydroxid titrated with double normal sulphuric acid, using litmus as indicator. By this method the formaldehyde is oxydized to formic acid, which is neutralized by the sodium hydroxid present.

While samples Nos. 65, 66, 67, and 26 I & W all contain less than 40 per cent formaldehyde by weight they contain 40 per cent or over by volume, so that the manufacturers are perfectly correct in claiming that their goods contain as much as 40 per cent of formaldehyde.

PETROLEUMS.

The results obtained by fractionally distilling four samples of crude petroleum are given in the following table:

TABLE	XVI	Composition of	f crude	vetroleum.
-------	-----	----------------	---------	------------

Temperature.	Character of distillate obtained.	Sample No. 275.b	Sample No. 276.c	Sample No. 277.d	Sample No. 278,¢
Below 50° C		Per cent.	Per cent.	Per cent.	Per cent.
50°-70° C	Petroleum ether				2. 0 1. 0
Below-80° C 80°-120° C 90°-120° C	Light naphtha Heavy naphtha Naphtha	1.49 4.35	0.14 1.63	1.89 9.26	
120°-150° C 150°-200° C 200°-250° C 250°-300° C	Benzine Light burning oils Heavy burning oils	5. 03 7. 64 13. 54	3. 82 13. 48 12. 03	9. 04 10. 73 15. 80	5.3 13.7 12.4 17.7
300°-315° C	Illuminating oilsdo	68.70	68, 62	45, 60 7, 70	23.93 16.93

α Zeit für anal. Chemie, 1897, p. 19. b Specific gravity, 35° Baumé.

c Specific gravity, 34° Baumé. d Specific gravity, 45° Baumé.

MISCELLANEOUS SOLID INSECTICIDES AND FUNGICIDES.

The analyses of a number of solid insecticides and fungicides that do not come under any of the previous headings are given in the following tables:

SULPHURS.

Table XVII.—Composition of sulphurs.

Constituents.	Sample No. 35.	Sample No. 36.
Soluble sulphur Insoluble sulphur Lime Foreign matter	Per cent. 99, 960 .144 .000 .196	Per cent. 98.666 .576 Trace .76
Total	100.300	100.000

The total for sample No. 35 is given by the chemist who made the analysis as 100 per cent, but all the constituents added together amount to 100.3 per cent, as above.

PINE TAR INSECTICIDES.

[No. 73.]

This is an insecticide obtained from the destructive distillation of pine knots. It readily mixes with water and is slightly sticky and greasy.

BUG DEATHS.

Table XVIII.—Composition of bug deaths.

Constituents.	Sample No. 93.	Sample No. 223.
	Per cent.	Per cent.
loisture	0.03	0.
ead oxid	1, 58	2.
ine oxid	78.86	86.
erric oxid and alumina	3. 80	5.
ilica		2.
oss on ignition		2.
oss on ignition hosphorie acid itrogen		
itrogen		
Potash		

POTASSIUM CYANID.

[Nos, 23 and 1170 I & W.]

Sample No. 23 contained 53.12 per cent of potassium cyanid calculated from the cyanogen. The remainder of the sample appeared to consist principally of moisture and some potassium carbonate and chlorid. The method of analysis used was that described in Bulletin 68, Bureau of Chemistry.

Composition of potassium cyanid.

[No. 1170, I & W.]	
	Per cent.
Cyanogen	 . 40, 49
Potassium	 . 21.48
Sodium	 23, 74
Chlorin	 8.98
Carbonic acid radical	 . 1.98
Moisture	
Undetermined	 2.51
	100.00
	100,00

The above analyses show that both samples of potassium cyanid are very impure, although the second sample contains just about the amount of cyanogen demanded by theory.

NONPOISONOUS POTATO-BUG DESTROYER.

Composition of nonpoisonous potato-buy destroyer.

[No. 137.]	
· ·	Per cent.
Moisture	0.00
Nitrogen	08
Ash	79,85
Potassium oxid	00
Calcium oxid	68.20
Magnesium oxid	7.29
Ferric oxid and alumina	1.38
Phosphorus pentoxid	Trace.
Insoluble before calcination	7.29
Insoluble after calcination	1.50

As this analysis is not complete, no opinion as to the substances of which the mixture is composed can be expressed.

NONPOISONOUS INSECT EXTERMINATOR.

[No. 151.]

This sample contained 78.12 per cent of lime and 3.61 per cent crude carbolic acid, determined as ether extract. It had an odor of carbolic acid and appeared to be partly composed of carbolate of lime.

X. O. DUST.

[No. 176.]

Composition of X. O. Dust.

	Per cent.
Moisture	6. 98
Organic and volatile matter	53. 69
Carbon dioxid	
Carbolic acid	Trace.

Mineral matter	39. 33
Sand	14.96
Ferric oxid and alumina	2.47
Calcium oxid	17. 10
Magnesium oxid	2.09
Phosphorus pentoxid	
Potassium oxid	

This mixture appears to be partly composed of carbonate of calcium with an excess of lime. No opinion as to the character of the organic and volatile matter can be based upon the data given, exclusive of the carbolic acid already mentioned.

ELECTRIC VERMIN EXTERMINATOR.

[No. 197.]

This insecticide is chiefly a mixture of calcium carbonate and calcium hydroxid with a very small amount of organic matter. Upon examination in this laboratory the organic matter present was found to consist of carbolic acid.

PAR' OIDIUM.

[No. 281.]

From the analyst's report this product would appear to consist of silicate of iron mixed with calcium sulphate. The analysis does not show the presence of sulphur and tobacco, both of which were found on examining a sample of the product in this laboratory.

ANTINONNIN.

[No. 282.]

A qualitative test of this substance showed that it contains dinitrocresols, together with soap and glycerin.

MISCELLANEOUS LIQUID INSECTICIDES AND FUNGICIDES.

BOLLENE INSECTICIDE.

Composition of samples Nos. 305, 644, 645.

[No. 305.]

	Per	r cent.
Carbon bisulphid		47.8
Mineral and essential oils		12.2
Water and residue		40.0

This analysis was made in accordance with the Texas law (see p. 61) regulating the sale of Paris Green and other insecticides and is good only until May 1, 1903.

[Nos, 644 and 645 (1 & W).]

Of the two samples of this product received at the Bureau of Chemistry one consisted of a black oil (644), while the other (645) separated into two layers, one a black oil and the other a white milky emulsion. The first sample was fractionally distilled as it stood, and the second sample was separated into its two parts and each part subjected to fractional distillation.

No. 644 (I & W), results of distillation test.

			Pe
Below 95° C		 	11
95°-108° C		 	
108°−130° C		 	
130°−160° C		 	
160°−190° C		 	
190°−220° C		 	
260°-300° C		 	e
350° C. +		 	
Loss and residu	ıe	 	

a Practically all carbon bisulphid.

No. 645 (I & W), results of distillation test.

WHITE EMULSION, 81.28 PER CENT.

	Per cent.
Below 95° C	a1.24
95°-108° C	b 76, 10
108°-130° C	c 13, 47
Loss and residue	9. 19

BLACK OIL, 18.72 PER CENT.

	Per cent.
Below 40° C	. 443.48
40°-65° C	. d 22.00
65°-95° C	. d 5, 53
95°-108° C	. e.50
108°-130° C	. e 4.84
130°-160° C	. f 8. 21
160° C. +	. f 6.53
Residue and loss	. 8.91

a Odor of earbon bisulphid.

b Penetrating odor of some essential oil.

c Odor of petroleum products with paraffin which crystallizes out in the higher fractions.

b Mostly water with solid particles of paraffin.

c Nearly all paraffin mixed with water.

d Nearly all carbon bisulphid.

e Some high-smelling essential oil present.

f Mineral oils with odor of petroleum products.

It is evident from the analyses of the two samples of bollene insecticide made in the laboratory that the first one (644 I & W) consists of about 50 per cent carbon bisulphid and 50 per cent crude petroleum with a very small quantity of some essential oil having a strong odor, while the second (645 I & W) consists of about 62 per cent water, 14 per cent carbon bisulphid, and 23 per cent petroleum, with a little essential oil and a small quantity of soap. Whether or not a compound of this composition, which is evidently very variable, would kill the bollworm is a matter for experimentation.

The results of the analyses of these two samples were forwarded to the manufacturer, who in reply said: "This analysis is not at all correct, and since the analysis was made I have perfected my preparation." Another letter was then sent, asking in what particular the analysis was unsatisfactory, but to this no reply was received. The analyses of these samples was repeated, but the same results were obtained. Besides this it is found that the analyses made in this laboratory gave very much the same results as those made by the Texas authorities from a qualitative standpoint. Unless the results obtained by the Texas authorities and by this laboratory are approximately correct the manufacturer is selling all of his goods under a false label, since the Texas law requires that all insecticides sold in the State shall comply with the official analysis on every package.

EXCELSIOR INSECTICIDE AND WOOD PRESERVER, RED CEDAR BRAND.

[No. 63, I & W.]

Results of distillation test.

	Per cent.
(1) 80°-110° C	a 1.01
(2) 110°-180° C	
(3) 180°–230° C	b 7.42
(4) 230°-270° C	b 21.91
(5) 270°-300° C	16.50
(6) 300°-330° C	19.11
(7) 330°-355° C	15, 84
(8) Residue and loss	17, 90

a Light oils with odor of benzene, toluene, etc.

Fractions 4, 5, 6, and 7 are heavier than water. At about 340° C. the distillate begins to solidify in the condenser, and at 355° C, the condenser is so stopped up with naphthalene that it is necessary to cease distilling. This appears to be a sample of coal-tar creosote, and when properly used is of value both as an insecticide and a wood preserver.

b Give tests for phenol and cresols.

FLY PREVENTATIVE.

[No. 202, I & W.]

Results of distillation test.

		Per	cent.
(1)	80°-160° C		$a _{8}$
(2)	160°-215° C		b 7
(3)	215°-270° C		€ 16
	270°-350° C		
(5)	350° C		d33
(6)	Residue and loss		e 17

 $[\]boldsymbol{a}$ Light oil giving test for phenol and cresols,

Fractions 1, 2, 3, 4, and 5 are all lighter than water. From the above analysis this sample appears to be a mixture of coal-tar creosote and petroleum. These substances are often used as fly preventatives.

KARBO KRESOLATE.

[No. 916, I & W.]

Results of distillation test.

	Per cent.
(1) 85°-105° C	a 26. 4
(2) 105°-180° C	
(3) 180°-220° C	
(4) 220°–270° C	
(5) 270°–285° C	e8.8
(6) 285°-312° C	14.7
(7) Residue and loss	28.7

a Light oil and water mixed; smell of carbon bisulphid and benzine.

Fractions 3 and 4 contain solid naphthalene. A small amount of alkali is also present. From the distillation test it would appear that this sample consists of coal-tar creosote which has been boiled with resin and an alkali, or with resin, fatty matter, and alkali, to cause it to form an emulsion with water. From the size, appearance, and smell of the first fraction it also appears that some carbon bisulphid has been added. It is well known that creosote emulsions have given good results both as disinfectants and sheep dips. This preparation, therefore, would very likely be of value for the purposes intended.

b Heavier oil giving test for phenol and cresols.

e Thick oils.

d Thick oil with solid particles.

e Solid when cooling.

b Light oil, test for phenol and cresols.

coil, heavier than water, test for phenol and cresols.

d Oil heavier than water.

e Thick oil; pungent odor.

PROFESSOR DE GRAFF'S CARPET BUG DESTROYER.

Composition of Professor De Graff's Carpet Bug Destroyer.

Constituents.	Sample No. 122.	Sample No. 129.
Moisture	Per cent. 95, 811 4, 189	Per cent. 95, 81
Mercury Chlorin Sulphur trioxid Alumina Potassium oxid	. 782 . 265 . 484 . 904 . 267	.78 .27 .48 90

These two analyses are given in two different reports of the Massachusetts State Experiment Station, but were very likely made upon the same samples, since they are identical.

The active poisonous principle of this solution is evidently mercuric chlorid, which is often used for the destruction of various insects which infest houses.

III. STATE LAWS GOVERNING THE COMPOSITION AND SALE OF INSECTICIDES.

A circular letter was sent to the various experiment stations asking that they forward copies of the laws in their respective States relative to the composition and sale of Paris Green and other insecticides. In cases where the experiment stations could not furnish the information a similar request was made of the secretary of state. It is believed that the following compilation of State laws is a complete résumé of all such laws passed in the United States.

CALIFORNIA.

AN ACT to prevent fraud in the sale of Paris Green used as an insecticide.

The people of the State of California, represented in senate and assembly, do enact as follows: It shall be the duty of each and every manufacturer of Paris Green (commercial aceto-arsenite of copper) to be used as an insecticide within this State, and of every dealer in original packages of said Paris Green manufactured outside of this State, before the said Paris Green is offered or exposed for sale, or sold within this State as an insecticide, to submit to the director of the California experiment station, at Berkeley, samples of said Paris Green, and a written or printed statement setting forth: First, the brand of said Paris Green to be sold, the number of pounds contained in each package in which it is put on the market for sale, the name or names of the manufacturers and the place of manufacturing the same; second, the statement shall set forth the amount of combined arsenic which the said Paris Green contains, and the statement so furnished shall be considered as constituting a guarantee to the purchaser that every package of such Paris Green contains not less than the amount of combined arsenic set forth in the statement.

Sec. 2. Every purchaser of said Paris Green in original packages which is manufactured outside of this State, who intends to sell or expose the same for sale, and every manufacturer of said Paris Green within this State, shall, after filing the statement above provided for with the director of the California agricultural station, at Berkeley, receive from the said director a certificate stating that he has complied with the foregoing statements, which certificate shall be furnished without charge therefor. Said certificate when furnished shall authorize the party when receiving the same to deal in this State in the said Paris Green. Any person who fails to comply with the terms of section 1 of this act shall not be entitled to such certificate and shall not be entitled to deal in said Paris Green within this State. Nothing in this section shall be construed as applying to retail dealers selling said Paris Green which has already been labeled and guaranteed.

SEC. 3. Paris Green, when sold, offered, or exposed for sale as an insecticide in this State shall contain at least 50 per centum of arsenious oxide and shall not contain more than 4 per centum of the same in the uncombined state.

SEC. 4. The director of the California State agricultural station, at Berkeley, shall examine or cause to be examined different brands of Paris Green sold, offered, or exposed for sale within this State, and cause samples of the same to be analyzed, and

shall report results of analysis forthwith to the secretary of the State board of horticulture and to the party or parties submitting such samples, and such report shall be final as regards the quality.

Sec. 5. Any person or persons, firm, association, company, or corporation violating any of the provisions of this act, and any person who shall sell any package of Paris Green, or any part thereof, which has not been labeled as herein provided, shall be guilty of a misdemeanor, and shall be fined not less than \$50 nor more than \$200, together with cost of the suit, in an action caused to be brought by the State board of horticulture, through its secretary, in the name of the people of the State of California.

Sec. 6. The attorney-general of the State of California is charged with the prosecution of all such suits.

Sec. 7. This act will take effect immediately.—Became a law by constitutional limitation February 28, 1901.

LOUISIANA.

Act No. 131 of the general assembly of the State of Louisiana.

Be it enacted by the general assembly of the State of Louisiana, That the bureau of agriculture shall be charged with the duties of regulating the sale and purity of Paris Green as an insecticide in this State.

Sec. 2. That it shall be the duty of any manufacturer or dealer in original packages of Paris Green before the same is offered for sale in this State, to submit to the commissioner of agriculture a written or printed statement setting forth: First. The brand of Paris Green to be sold, the number of pounds contained in each package in which it is to be put upon the market for sale, and the name or names of the manufacturers and the place of manufacture. Second. The statement setting forth the amount of arsenic which they are willing to guarantee the said Paris Green to contain, and the statement so furnished shall be considered as constituting a guarantee to the purchaser that every package of said Paris Green contained not less than the amount of arsenic set forth in the statement.

Sec. 3. That every purchaser proposing to deal in Paris Green shall, after filing the statement above provided for with the commissioner of agriculture, receive from the said commissioner of agriculture a certificate stating that he has complied with the foregoing section, which certificate shall be furnished by the commissioner without any charge therefor; that the said certificate when furnished shall authorize the party receiving the same to deal in this State in Paris Green; that no person who has failed to file the statement aforesaid and to receive the certificate of authority aforesaid shall be authorized to deal in this State in Paris Green, and any person so dealing in this State without having filed the aforesaid statement and received the certificate aforesaid shall be liable for each violation to a fine not exceeding \$250, which fine shall be recoverable before any court of competent jurisdiction at the suit of the commissioner of agriculture or of any citizen, and shall be disposed of as hereinafter provided; provided, further, that nothing in this section shall be construed as preventing the sale by retail dealers throughout the State of Paris Green which has already been guaranteed and labeled as provided for in this act.

Sec. 4. That it shall be the duty of the board of agriculture, or its commissioner, at the opening of each season, to issue and distribute circulars setting forth the brands of Paris Green, their percentages of arsenic as claimed by the dealers, and to more particularly describe them they shall be separated into two classes, viz: First, those brands containing 50 per cent or more of arsenic shall be classed as "strictly pure," and, second, all falling below this percentage shall be classed as "impure."

Sec. 5. That it shall be the duty of the commissioner of agriculture to cause to be prepared labels of suitable material, fitted to be attached to packages of Paris Green and to have printed thereon "guaranteed," with a blank space into which there may

be stamped by the commissioner of agriculture the words "strictly pure," or "impure," as the guarantee may require; also the year or season in which it is to be used and a facsimile of the signature of said commissioner. The said labels shall be furnished by the said commissioner, to any dealer in Paris Green who shall have complied with the foregoing provisions of this act, upon the payment by said dealer to said commissioner of fifty cents for a sufficient number to label one hundred pounds of Paris Green.

Sec. 6. That it shall be the duty of every person, before offering for sale any Paris Green as an insecticide in this State, to attach or cause to be attached to each package one of the labels hereinbefore described, designating the quantity of Paris Green in the package to which it is attached. Any person who shall sell any packages of Paris Green, or any part thereof, which has not been labeled as herein provided for shall be guilty of a misdemeanor, and on conviction thereof shall be fined in the sum of one hundred dollars for each omission, which penalty may be sued for either by the Commissioner of Agriculture or any person for the uses hereinafter declared. Any person who shall counterfeit or use a counterfeit label prescribed by this act, or who shall use them a second time, shall be guilty of a misdemeanor, and on conviction thereof shall be fined in a sum not exceeding two hundred and fifty dollars, one-half of which shall be paid to the informer, which may be doubled or trebled at each second or third conviction, and so on progressively for subsequent convictions.

SEC. 7. That it shall be the duty of every person who sells a package of Paris Green, upon the request of the purchaser, to draw from the same and in the presence of the purchaser or his agent a fair or correct sample and to have the same securely enclosed and sealed and sent to the Commissioner of Agriculture for analysis by the official chemist of the State, and if upon analysis the said Paris Green shall be found below the guarantee given to the Commissioner of Agriculture and printed on the package, then the said seller shall be liable to the said purchaser for all damage accruing from said difference, recoverable in any court of competent jurisdiction in the State.

Sec. 8. That the copy of the official chemist's analysis of any Paris Green certified to by him shall be admissible as evidence in any court of the State on trial of any issue involving the merits of said Paris Green.

SEC. 9. That the Bureau of Agriculture shall adopt needful rules and regulations providing for the collecting of money arising from the sale of labels, or for any fines imposed under this act, and shall deposit the same with the Treasurer of the State.

NEW YORK.

AN ACT to amend the agricultural law, to prevent fraud in the sale of Paris Green.

The People of the State of New York, represented in Senate and Assembly, do enact as follows: Chapter three hundred and thirty-eight of the laws of eighteen hundred and ninety-three entitled "An act in relation to agriculture, constituting articles one, two, three, four, and five, of chapter thirty-three of the general laws," is hereby amended by adding a new article to be known as article eight and to read as follows:

SEC. 110. State manufacturer and the dealer in original packages to file certificate with commissioner of agriculture. After the passage of this act it shall be the duty of each and every manufacturer of Paris green within this state, and of every dealer in original packages of Paris green manufactured outside of this state, before the said Paris green is offered or exposed for sale or sold within this state, to submit to the commissioner of agriculture a written or printed statement setting forth, first, the brands of Paris green to be sold, the number of pounds contained in each package in which it is put upon the market for sale, the name or names of the manufacturers and the place of manufacturing the same; second, the statement shall set forth the amount of arsenic which the said Paris green contains, and the statement so furnished

shall be considered as constituting a guarantee to the purchaser that every package of such Paris green contains not less than the amount of arsenic set forth in the statement.

Sec. 111. Certificate to be given by the commissioner of agriculture to state manufacturer and dealer in original packages. Every purchaser of Paris green in original packages, which is manufactured outside of this state, who intends to sell or expose the same for sale, and every manufacturer of Paris green within this State, shall, after filing the statement above provided for with the commissioner of agriculture, receive from the said commissioner of agriculture a certificate stating that he has complied with the foregoing statement, which certificate shall be furnished without any charge therefor; said certificate when furnished shall authorize the party receiving the same to deal in this state in Paris green. Any person who fails to file the statement aforesaid shall not be entitled to such certificate and shall not be entitled to deal in Paris green within this State; nothing in this section shall be construed as applying to retail dealers.

Sec. 112. Composition of Paris green or analogous products. Paris green or any product analogous to it, when sold, offered or exposed for sale, as such, in this state, shall contain at least fifty per centum of arsenious oxide. (See Amendment.)

SEC. 113. Paris green to be analyzed at experiment station. The director of the New York State agricultural experiment station, at Geneva, shall, under the direction of the commissioner of agriculture, examine, or cause to be examined, the different brands of Paris green sold, offered, or exposed for sale within the State, and cause samples of the same to be analyzed, and shall report the result of the analysis forthwith to the commissioner of agriculture.

Sec. 114. Penalty for violations. Any person or persons, firm, association, company, or corporation violating any of the provisions of this act shall be guilty of a misdemeanor, and shall be fined not less than fifty dollars nor more than two hundred dollars; and in addition thereto shall forfeit and pay unto the people of the state of New York the sum of one hundred dollars, together with the costs of the suit in an action caused to be brought by the commissioner of agriculture in the name of the people of the state of New York, as provided by section eight of the agricultural law.

2. This act shall take effect immediately. Became a law March 23, 1898, by the approval of the Governor. (Laws of New York, ch. 113.)

AMENDMENT TO PARIS-GREEN LAW.

In accordance with the suggestion made by the New York State Experiment Station in 1900, that portion of the Paris Green law which related to the definition of Paris Green was changed. The essential portion of the amended law embodying this change is as follows:

Sec. 112. Composition of Paris green or analogous products. Paris green, or any product analogous to it, when sold, offered, or exposed for sale as such in this state, shall comply with the following requirements:

First. It shall contain arsenic in combination with copper, equivalent to not less than fifty per centum arsenious oxide.

Second. It shall not contain arsenic in water-soluble forms equivalent to more than three and one-half per centum of arsenious oxide.

OREGON.

Only those sections of the act named below which deal directly with the subject of insecticides are given.

AN ACT to protect the fruit and hop industry of Oregon.

Sec. 3. It shall be unlawful for any person, firm or corporation doing business in the State of Oregon to sell paris green, arsenic, london purple, sulphur or any spray material or compound for spraying purposes, in quantities exceeding one pound, without providing with each package sold a certificate duly signed by the seller thereof guaranteeing the quality and per cent of purity of said materials.

Sec. 4. Any person, firm or corporation selling any of the above materials which do not conform with the certificate furnished therewith shall be deemed guilty of a misdemeanor, and upon conviction thereof shall be subject to a fine of not less than

twenty-five (\$25) dollars nor more than one hundred (\$100) dollars.

Sec. 8. It shall be the duty of the Commissioner of the State Board of Horticulture of the district in which a violation of this act occurs to present the evidence of the case to the district attorney, whose duty it shall be to prosecute any person guilty of a violation of this act, which prosecution may be brought in any of the justice courts of this state.

Sec. 9. Inasmuch as the horticultural interests of the state demand immediate attention, this act shall be in full force and effect from and after its approval by the Governor. Approved by the Governor, February 17, 1899.

TEXAS.

AN ACT for the better protection of the farmer in the purchase of commercial fertilizers and commercial poisons used for destroying bollworms and other pests.

Be it enacted by the Legislature of the State of Texas: Before any commercial fertilizer or commercial poison, or any chemical or mixture used as a commercial fertilizer or commercial poison, such as London purple, arsenic, Paris green, or any poison used for the purpose of destroying the boll worm or other pests, is sold or offered for sale in this State, the manufacturer, agent, importer or party who sells or offers it for sale within this State shall deposit with the professor of chemistry of the Agricultural and Mechanical College a sealed tin can, bottle or jar containing not less than one pound of the fertilizer or commercial poison offered for sale; with an affidavit that it is a fair sample taken from several barrels, boxes, sacks, or from quantities in larger bulk, of the article thus to be sold or offered for sale; provided the unmixed substances, cotton seed meal, land plaster, salt, ashes, lime, green sand marl, uncrushed bones and animal excrements shall be exempt from the operation of the law.

SEC. 2. The manufacturer, importer, vender or agent of any commercial fertilizer or commercial poison, as referred to in Section 1 of this chapter, shall pay annually to the treasurer of the Agricultural and Mechanical College an analysis fee of fifteen dollars for each and every fertilizer or commercial poison sold, exposed or offered for sale within this State. Such payment shall be made at the time the sample of fertilizer or commercial poison is submitted to the professor of chemistry for analysis.

Sec. 3. After the analysis fee has been paid, as provided for in Section 2 of this chapter, it shall be the duty of the professor of chemistry of the Agricultural and Mechanical College to analyze or to have analyzed under his direction, any sample of a commercial fertilizer or commercial poison, in accordance with the requirements of the foregoing sections of the law. The professor of chemistry shall print the

result of such analysis in the form of a label, which shall set forth the name of the manufacturer, the brand of such fertilizer or commercial poison, and the essential ingredients contained in such fertilizer or commercial poison, viz:

- 1. Available nitrogen and its equivalent in ammonia.
- 2. Soluble phosphoric acid; total available phosphoric acid.
- 3. Reverted phosphoric acid.
- 4. Total phosphoric acid.
- 5. Potash soluble in water.

This, however, shall not preclude the professor of chemistry from setting forth any other ingredient which the fertilizer may contain. And he shall place upon each label the money value of such fertilizer or commercial poison, computed from its composition as he may determine. He shall furnish such labels in quantities of 500 or multiple thereof at a cost of one dollar per hundred; the money to be paid directly to the treasurer of the Agricultural and Mechanical College.

Sec. 4. Every box, barrel, keg, or other package or quantity of commercial fertilizer or commercial poison (within the limitations of Section 1 of this law), in any shape or form, sold or offered for sale in this State, shall have attached to it in a conspicuous place the label as provided for in Section 3 of this law, with the signature of the professor of chemistry attached.

Sec. 5. The professor of chemistry, or any duly authorized agent of his, is hereby authorized to select from any package of commercial fertilizer or commercial poison sold, or exposed for sale in this State, a quantity not to exceed two pounds, for a sample to be used for the purpose of an official analysis and for comparison with the sample furnished by the manufacturer, agent, or vendor, for official analysis.

Sec. 6. Any manufacturer, agent, or vendor of any commercial fertilizer or commercial poison who shall offer or expose for sale any such fertilizer or commercial poison without having previously complied with the provisions of this chapter shall be fined not less than fifty and not more than five hundred dollars for each violation or evasion of this law.

Sec. 7. Any agriculturist or farmer, a purchaser of any commercial fertilizer or commercial poison in this State, may take a sample of the same under rules and regulations to be prescribed by the professor of chemistry of the Agricultural and Mechanical College and forward the same to him for analysis, which analysis shall be made free of charge.

Sec. 8. The revenues accruing from analysis fees and sale of labels, as provided for in this chapter, shall be expended by the board of directors of the Agricultural and Mechanical College for the maintenance of the chemical department and for such other purposes as they may determine.

Sec. 9. A copy of the official analysis of any fertilizer or commercial poison, or chemical certified to by the professor of chemistry, shall be admissible as evidence in any court of this State on the trial of any issue involving the merits of said fertilizer or commercial poison.

Sec. 10. Whereas the fact that there is no law authorizing the analysis of commercial fertilizers and commercial poisons, or the testing of adulterated poisons when used for the purpose of destroying the boll worm and other pests, and the better protection of the farmer, creates an emergency, and an imperative public necessity that the constitutional rule requiring bills to be read on three several days be suspended and that this act take effect and be in force from and after its passage, and said rule is so suspended and it is so enacted. Passed by the Twenty-sixth Legislature of Texas, 1899.

WASHINGTON.

AN ACT to provide against the adulteration of Paris Green and other compounds used for spraying trees and plants,

Be it enacted by the Legislature of the State of Washington, That it shall hereafter be unlawful for any person, firm, or corporation doing business in the State of Washington to sell or offer for sale adulterated or low grade Paris green, arsenic, London purple, sulphur, or any spray material or compound for spraying purposes.

Sec. 2. For the purposes of this act Paris green shall contain not less than fifty per cent of arsenic trioxide in combination, and not more than four per cent of water-soluble arsenic trioxide, and commercial arsenic shall contain not less than ninety-

six (96) per cent of arsenic trioxide.

Sec. 3. Any person, firm, or corporation violating any of the provisions of this act shall be deemed guilty of a misdemeanor, and upon conviction thereof shall be punished by a fine of not less than twenty-five dollars nor more than one hundred dollars. All fines imposed for violation of the provisions of this act shall be paid to the treasurer of the county wherein the violation was committed and placed to the general fund of such county.

Sec. 4. That the State Commissioner of Horticulture and the county fruit inspectors under his supervision be charged with the enforcement of this act, with the assistance

of the prosecuting attorney.

SEC. 5. It shall be the duty of the chemist of the State Agricultural Experiment Station to correctly analyze, without extra compensation and without extra charge to the state other than the necessary expenses, all substances and compounds used or offered for sale for spraying trees and plants, that the State Commissioner of Horticulture may send for analysis, and report to him without unnecessary delay the result of any analysis so made; any such chemist shall assist him in prosecuting violations of the law, by giving testimony, expert or otherwise. Approved by the Governor February 26, 1901.





RETURN TO the circulation desk of any University of California Library or to the

NORTHERN REGIONAL LIBRARY FACILITY Bldg. 400, Richmond Field Station University of California Richmond, CA 94804-4698

ALL BOOKS MAY BE RECALLED AFTER 7 DAYS

- 2-month loans may be renewed by calling (510) 642-6753
- 1-year loans may be recharged by bringing books to NRLF
- Renewals and recharges may be made 4 days prior to due date

DUE AS STAMPED BELOW

MAY 29 2003

JUL 29 2003

ELEY

DD20 15M 4-02

(P)s

